

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: August 13, 2005, 01:08:33 ; Search time 725.234 Seconds
(without alignments)
1536.704 Million cell updates/sec

Title: US-10-673-854-1

Perfect score: 23
Sequence: 1 catgtatttgatgggagatagagg 23

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl:.*
1: gb_ba.*
2: gb_htg.*
3: gb_in.*
4: gb_on.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_sts.*
12: gb_sy.*
13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	23	100.0	68726	9 AC016681	AC016681 Homo sapi
2	23	100.0	100545	9 AL590379	AL590379 Human DNA
3	23	100.0	129357	9 AL590492	AL590492 Human DNA
4	19.8	86.1	285888	2 AC107131	AC107131 Rattus no
5	18.8	81.7	3191	10 BC070398	BC070398 Mus muscu
6	18.8	81.7	5634	9 AB023160	AB023160 Homo sapi
7	18.8	81.7	8225	10 MMU420922	AJ420922 Mus muscu
8	18.8	81.7	20838	9 AC117508	AC117508 Homo sapi
9	18.8	81.7	25901	9 AC139617	AC139617 Homo sapi
10	18.8	81.7	134296	9 HS352H24	AL449215 Homo sapi
11	18.8	81.7	151700	9 AC133528	AC133528 Homo sapi
12	18.8	81.7	163947	2 AC102466	AC102466 Mus muscu
13	18.8	81.7	167797	2 AC151841	AC151841 Mus muscu
14	18.8	81.7	171446	2 AC118744	AC118744 Mus muscu
15	18.8	81.7	175280	10 AC113270	AC113270 Mus muscu
16	18.8	81.7	176364	10 AC113126	AC113126 Mus muscu
17	18.8	81.7	185571	9 AC133781	AC133781 Homo sapi
18	18.8	81.7	191154	2 AC132150	AC132150 Homo sapi
19	18.8	81.7	193634	2 AC068495	AC068495 Mus muscu

C 20	18.8	81.7	193894	9 AC114730	AC114730 Homo sapi
C 21	18.8	81.7	200711	2 AC133960	AC133960 Homo sapi
C 22	18.8	81.7	229639	5 AC140947	AC140947 Gallus ga
C 23	18.8	81.7	238704	2 BX890628	BX890628 Mus muscu
C 24	18.8	81.7	258520	10 AL662812	AL662812 Mouse DNA
C 25	18.4	80.0	450	8 ATSRG3PRT	X98376 A.thaliana
C 26	18.4	80.0	1117	8 AY114073	AY114073 Arabidops
C 27	18.4	80.0	1292	8 AY089160	AY089160 Arabidops
C 28	18.4	80.0	1307	8 AY072127	AY072127 Arabidops
C 29	18.4	80.0	1501	8 BT003327	BT003327 Arabidops
C 30	18.4	80.0	3050	6 CQ849723	CQ849723 Sequence
C 31	18.4	80.0	3050	9 AK126763	AK126763 Homo sapi
C 32	18.4	80.0	74188	2 AC101144	AC101144 Mus muscu
C 33	18.4	80.0	103960	8 ATAC011664	ATAC011664 Arabidops
C 34	18.4	80.0	110000	2 AC091341	AC091341 Rattus no
C 35	18.4	80.0	110000	2 AC114711_3	Continuation (4 of
C 36	18.4	80.0	158599	9 AC108066	AC108066 Homo sapi
C 37	18.4	80.0	163277	2 AC074259	AC074259 Trypanoso
C 38	18.4	80.0	163463	10 AC134536	AC134536 Mus muscu
C 39	18.4	80.0	173268	2 AC110225	AC110225 Mus muscu
C 40	18.4	80.0	181683	10 AC122180	AC122180 Mus muscu
C 41	18.4	80.0	221631	9 AC010867	AC010867 Homo sapi
C 42	18.4	80.0	223728	2 AC135443	AC135443 Rattus no
C 43	18.4	80.0	240364	2 AC114875	AC114875 Rattus no
C 44	18.4	80.0	256385	2 AC128721	AC128721 Rattus no
C 45	18.4	80.0	306803	1 AE017161	AE017161 Prochloro

ALIGNMENTS

RESULT 1	AC016681	68726 bp	DNA	linear	PRI 30-SEP-2000
LOCUS	Homo sapiens BAC clone RP11-62H15	from Y, complete sequence.			
DEFINITION	AC016681				
ACCESSION	AC016681.2	GI:7321924			
VERSION	HTG.				
KEYWORDS	Homo sapiens (human)				
SOURCE	Homo sapiens				
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
REFERENCE	1 (bases 1 to 68726)				
AUTHORS	Sulston,J.E. and Waterston,R.				
TITLE	Toward a complete human genome sequence				
JOURNAL	Genome Res. 8 (11), 1097-1108 (1998)				
MEDLINE	99063792				
PUBMED	9847074				
REFERENCE	2 (bases 1 to 68726)				
AUTHORS	Joshu,C., Strommatt,C. and Wedgeworth,P.				
TITLE	The sequence of Homo sapiens BAC clone RP11-62H15				
JOURNAL	Unpublished				
REFERENCE	3 (bases 1 to 68726)				
AUTHORS	Waterston,R.H.				
TITLE	Direct Submission				
JOURNAL	Submitted (04-DEC-1999) Genome Sequencing Center, Washington				
REFERENCE	4 (bases 1 to 68726)				
AUTHORS	Waterston,R.H.				
TITLE	Direct Submission				
JOURNAL	Submitted (24-MAR-2000) Genome Sequencing Center, Washington				
REFERENCE	5 (bases 1 to 68726)				
AUTHORS	Waterston,R.H.				
TITLE	Direct Submission				
JOURNAL	Submitted (04-APR-2000) Genome Sequencing Center, Washington				
REFERENCE	6 (bases 1 to 68726)				
AUTHORS	Waterston,R.H.				
TITLE	Direct Submission				

JOURNAL

Submitted (07-APR-2000) Genome Sequencing Center, Washington
University School of Medicine, 444 Forest Park Parkway, St. Louis,
MO 63108, USA
7 (bases 1 to 68726)

REFERENCE
AUTHORS

Waterston,R.

JOURNAL
TITLE

Direct Submission
Submitted (30-SEP-2000) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Mar 24, 2000 this sequence version replaced gi:6524399.

COMMENT

----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc>
Contact: sapiens@wustl.wustl.edu
----- Summary Statistics

Center project name: H_NH0062H15

NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
clone sections once, or longer because we provide a small overlap
between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate
chemistry, or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by sequence
from more than one subclone; and the assembly was confirmed by
restriction digest.

MAPPING INFORMATION:

The position of this clone was established as part of a
collaboration between the Human Chromosome Y Mapping Project
(Tomoko Kawaguchi, Helen Skaletsky, Laura G. Brown, Steve Rozen,
and David C. Page at the Whitehead Institute for Biomedical
Research, Cambridge MA) and the Washington University Genome
Sequencing Center, St. Louis MO.

SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male
donor, as described by Osoegawa,K., Moon,P.Y., Zhao,B., Frengen,E.,
Tateno,M., Catanesi,J.J. and de Jong,P.J. (1998) An improved
approach for construction of bacterial artificial chromosome
libraries. Genomics 51:1-8. The clone may be obtained either from
Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong
and coworkers at the Roswell Park Cancer Institute
(<http://bacpac.med.buffalo.edu>)
VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-362J16; the clone sequenced
to the right is RP11-218E11. Actual start of this clone is at base
position 1 of RP11-62H15.

FEATURES

source

1..68726
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="Y"
/map="Y"
/clone="RP11-62H15"
/clone_lib="RPCI-11"

repeat_region

1..352

repeat_region

357..651

repeat_region

676..769

repeat_region

1837..1873

repeat_region

2274..2301

repeat_region

2542..2572

repeat_region
3768..3973
/rpt_family="AT_rich"
repeat_region
4258..4660
/rpt_family="MaLR"
repeat_region
5234..5568
/rpt_family="MaLR"
repeat_region
6159..6713
/rpt_family="MER1_type"
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6718..6943
/rpt_family="L1"
repeat_region
6929..7694
/rpt_family="L1"
repeat_region
7709..7887
/rpt_family="L1"
repeat_region
7903..8247
/rpt_family="TA)n"
repeat_region
8264..9031
/rpt_family="L1"
repeat_region
9435..9583
/rpt_family="MIR"
repeat_region
9584..9738
/rpt_family="MIR"
repeat_region
10428..10482
/rpt_family="L2"
repeat_region
10574..11396
/rpt_family="L1"
repeat_region
11997..12024
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repeat_region
12040..13387
/rpt_family="L1"
repeat_region
13390..13846
/rpt_family="L1"
repeat_region
13837..14070
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repeat_region
14074..15649
/rpt_family="L1"
repeat_region
15648..16717
/rpt_family="L1"
repeat_region
16720..16789
/rpt_family="A-rich"
repeat_region
16827..17303
/rpt_family="L1"
repeat_region
17417..17575
/rpt_family="L2"
repeat_region
17725..17768
/rpt_family="AT_rich"
repeat_region
18047..18132
/rpt_family="L1"
repeat_region
18338..18648
/rpt_family="Alu"
repeat_region
19586..19620
/rpt_family="CA)n"
repeat_region
19704..19918
/rpt_family="MIR"
repeat_region
19925..19980
/rpt_family="ATG)n"
repeat_region
20305..20330
/rpt_family="CA)n"
repeat_region
20333..20671
/rpt_family="MER2_type"
repeat_region
21471..21557
/rpt_family="MaLR"
repeat_region
21661..22655
/rpt_family="L1"
repeat_region
22725..22971
/rpt_family="L1"
repeat_region
23441..23462
/rpt_family="AT_rich"
repeat_region
23982..24011
/rpt_family="AT_rich"
repeat_region
24167..24258
/rpt_family="L1"

repeat_region 24399..24463
 /rpt_family="AT_rich"
 repeat_region 24468..24576
 /rpt_family="L2"
 repeat_region 24754..25021
 /rpt_family="Retroviral"
 repeat_region 25115..25575
 /rpt_family="L2"
 repeat_region 25715..26221
 /rpt_family="MaLR"
 repeat_region 26501..26523
 /rpt_family="AT_rich"
 repeat_region 26567..26925
 /rpt_family="MaLR"

Query Match 100.0%; Score 23; DB 9; Length 68726;
 Best Local Similarity 100.0%; Pred. No. 2.9;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CATGTATTGATGGGATAGAGG 23
 Db 56609 CATGTATTGATGGGATAGAGG 56631
 |||||
 |||||

RESULT 2
 AL590379
 LOCUS
 DEFINITION Human DNA sequence from clone RP11-343H6 on chromosome X, complete sequence.
 ACCESSION AL590379.7 GI:29466482
 VERSION AL590379
 KEYWORDS HTG.
 SOURCE Homo sapiens (human)
 ORGANISM
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 100545)
 AUTHORS Howden, P.
 TITLE Direct Submission
 JOURNAL Submitted (01-APR-2003) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
 COMMENT On Apr 1, 2003 this sequence version replaced gi:28933281.
 ----- Genome Center
 Center: Wellcome Trust Sanger Institute
 Center code: SC
 Web site: <http://www.sanger.ac.uk>
 Contact: humquery@sanger.ac.uk

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.
 This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest, except on the rare occasion of the clone being a YAC.
 The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em., EMBL; Sw., SWISSPROT; Tr., TrEMBL; Wp., WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/ChrX>
 RP11-343H6 is from the library RPCI-11.2 constructed by the group

of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>
 VECTOR: pBACE3.6.

FEATURES
 source 1..100545
 Location/Qualifiers
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosome="X"
 /clone="RP11-343H6"
 /clone_lib="RPCI-11.2"

ORIGIN

Query Match 100.0%; Score 23; DB 9; Length 100545;
 Best Local Similarity 100.0%; Pred. No. 2.9;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CATGTATTGATGGGATAGAGG 23
 Db 67470 CATGTATTGATGGGATAGAGG 67492
 |||||
 |||||

RESULT 3
 AL590492
 LOCUS
 DEFINITION Human DNA sequence from clone RP11-88H5 on chromosome X, complete sequence.
 ACCESSION AL590492
 VERSION AL590492.7 GI:15072594
 KEYWORDS HTG.
 SOURCE Homo sapiens (human)
 ORGANISM
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 129357)
 AUTHORS Heath, P.
 TITLE Direct Submission
 JOURNAL Submitted (10-DEC-2001) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
 COMMENT On Aug 1, 2001 this sequence version replaced gi:1329489. During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.
 This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em., EMBL; Sw., SWISSPROT; Tr., TrEMBL; Wp., WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/ChrX>
 RP11-88H5 is from the library RPCI-11.1 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>
 VECTOR: pBACE3.6

IMPORTANT: This sequence is not the entire insert of clone RP11-88H5 it may be shorter because we sequence overlapping sections only once, except for a short overlap. The true left end of clone RP11-88H5 is at 1 in this sequence. The true left end of clone RP11-156J23 is at 129258 in this sequence.
 Location/Qualifiers

FEATURES

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/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosomes="X"
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/clone lib="RPC1-11.1"
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1933. .2582
/note="L1M4 repeat: matches 2318. .3033 of consensus"
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complement(2286. .2752)
/note="match: GSS: Em:AQ450883"
misc_feature
complement(2291. .2748)
/note="match: GSS: Em:AQ360394"
misc_feature
2846. .3095
/note="match: STS: Em:HSC28P10"
repeat_region
2303. .3363
/note="MLT1H repeat: matches 5. .546 of consensus"
misc_feature
4611. .5105
/note="match: GSS: Em:AQ827938"
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5147. .5317
/note="L1P4 repeat: matches 5423. .5594 of consensus"
repeat_region
5634. .6010
/note="THE1C repeat: matches 1. .371 of consensus"
repeat_region
6017. .6215
/note="MTC repeat: matches 1. .187 of consensus"
repeat_region
6369. .6505
/note="MTRC repeat: matches 254. .394 of consensus"
repeat_region
7390. .7767
/note="L1ME2 repeat: matches 4453. .4836 of consensus"
repeat_region
7776. .8020
/note="L1MC2 repeat: matches 6070. .6329 of consensus"
repeat_region
8028. .8148
/note="MER7A repeat: matches 445. .558 of consensus"
repeat_region
8174. .8227
/note="MER7A repeat: matches 412. .466 of consensus"
repeat_region
8228. .8523
/note="AluY repeat: matches 3. .297 of consensus"
repeat_region
8524. .8921
/note="MER7A repeat: matches 26. .412 of consensus"
repeat_region
8937. .9122
/note="L1MC1 repeat: matches 5870. .6072 of consensus"
repeat_region
9142. .9431
/note="AluY repeat: matches 1. .289 of consensus"
repeat_region
9432. .9818
/note="L1ME2 repeat: matches 4845. .5248 of consensus"
repeat_region
10083. .10417
/note="MLT1E repeat: matches 240. .568 of consensus"
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10418. .10564
/note="L1ME2 repeat: matches 5237. .5397 of consensus"
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10568. .11070
/note="MLT1D repeat: matches 1. .500 of consensus"
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11071. .11816
/note="L1ME2 repeat: matches 5391. .6153 of consensus"
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11825. .12014
/note="L1ME2 repeat: matches 1856. .2055 of consensus"
repeat_region
12146. .12258
/note="MLT1E repeat: matches 28. .136 of consensus"
repeat_region
12295. .12528
/note="MLT1E-internal repeat: matches 1114. .1335 of consensus"
repeat_region
12547. .12965
/note="MLT2CB repeat: matches 1. .433 of consensus"
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13623. .14146
/note="MER9 repeat: matches 1. .511 of consensus"
repeat_region
14225. .14517
/note="MLT1-INTERNAL repeat: matches 291. .591 of consensus"
repeat_region
16863. .17118
/note="L1ME3A repeat: matches 5386. .5638 of consensus"
repeat_region
17119. .17250
/note="L1PB1 repeat: matches 6024. .6155 of consensus"

repeat_region
17251. .17506
/note="L1MB3A repeat: matches 5638. .5903 of consensus"
repeat_region
17546. .17932
/note="L1M4 repeat: matches 251. .627 of consensus"
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18147. .18210
/note="L1M4 repeat: matches 616. .674 of consensus"
repeat_region
18184. .19717
/note="L1M4 repeat: matches 926. .2109 of consensus"
repeat_region
19782. .19970
/note="L1M4 repeat: matches 2123. .2308 of consensus"
repeat_region
19975. .20288
/note="AluO repeat: matches 12. .312 of consensus"
repeat_region
20314. .20637
/note="L1M4 repeat: matches 2316. .2663 of consensus"
repeat_region
20644. .21468
/note="L1P3 repeat: matches 15. .650 of consensus"
repeat_region
21469. .23040
/note="L1PA13 repeat: matches 4584. .6152 of consensus"
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/note="match: GSS: Em:B79974"
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23281. .23580
/note="AluO repeat: matches 1. .291 of consensus"
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23909. .24217
/note="AluSx repeat: matches 1. .309 of consensus"
repeat_region
24276. .24322
/note="MADL1 repeat: matches 33. .79 of consensus"
repeat_region
24326. .24482
/note="L2 repeat: matches 1992. .2147 of consensus"
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24990. .25140
/note="L1M4 repeat: matches 2404. .2560 of consensus"
repeat_region
25585. .25883
/note="L1 repeat: matches 3095. .3404 of consensus"
repeat_region
25903. .26200
/note="AluY5 repeat: matches 1. .296 of consensus"
repeat_region
26529. .26685
/note="FRAM repeat: matches -4. .152 of consensus"
repeat_region
26825. .26996
/note="MLT1J repeat: matches 320. .509 of consensus"
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27032. .27116
/note="HERV16 repeat: matches 2. .89 of consensus"
repeat_region
27137. .27444
/note="L1R16A repeat: matches 115. .426 of consensus"
repeat_region
28310. .28924
/note="L1M4 repeat: matches 5064. .5692 of consensus"
repeat_region
29682. .29969
/note="AluSx repeat: matches 1. .288 of consensus"
repeat_region
30022. .30136
/note="L1MB1 repeat: matches 6033. .6148 of consensus"
repeat_region
30157. .34776
/note="L1PA13 repeat: matches 648. .5299 of consensus"
repeat_region
34850. .35692
/note="L1PA13 repeat: matches -651. .215 of consensus"
repeat_region
35729. .36642
/note="L1 repeat: matches 3630. .4551 of consensus"
repeat_region
36652. .36809
/note="MLT1A1 repeat: matches 1. .160 of consensus"
repeat_region
36810. .38853
/note="L1P repeat: matches 3236. .5271 of consensus"
repeat_region
38877. .39968
/note="L1P repeat: matches 2318. .3413 of consensus"
repeat_region
40002. .40330
/note="L1P repeat: matches 2201. .2320 of consensus"
repeat_region
40118. .40327
/note="AluY repeat: matches 94. .303 of consensus"
repeat_region
40328. .41566
/note="L1P repeat: matches 2656. .3894 of consensus"
repeat_region
41571. .41865
/note="AluB repeat: matches 1. .288 of consensus"
repeat_region
41890. .41983
/note="L1 repeat: matches 3856. .3955 of consensus"
repeat_region
44982. .45187
/note="MLT1 repeat: matches 1. .205 of consensus"
repeat_region
45472. .45874
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/note="MLR11 repeat: matches 1. .402 of consensus"
46448. .46782
/note="MER58B repeat: matches 2. .341 of consensus"
47373. .46157
/note="L1 repeat: matches 4164. .4992 of consensus"

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```

Query Match      100.0%; Score 23; DB 9; Length 129357;
Best Local Similarity 100.0%; Pred. NO. 2.9;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 CATGTATTTCATGGGATAGG 23
Db 97822 CATGTATTTCATGGGATAGG 97844

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RESULT 4
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LOCUS      285888 bp      DNA      linear      HTG 22-SEP-2002
DEFINITION Rattus norvegicus clone CH230-38C17, *** SEQUENCING IN PROGRESS
***, 6 unordered pieces.
AC107131.4 GI:23267848
VERSION    HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.
KEYWORDS   Rattus norvegicus
SOURCE     Rattus norvegicus
ORGANISM   Mammalia; Eutheria; Chordata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
          Rattus.

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REFERENCE
AUTHORS    Muzny,D,Marie., Metzker,M, Lee., Abramson,S., Adams,C., Alder,J.,
          Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
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TITLE
JOURNAL
REFERENCE
AUTHORS
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AUTHORS
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JOURNAL
COMMENT

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Williams,G., Willson,R., Wlaczek,R., Wooden,H., Worley,K., Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V., Yu,P., Zhang,J., Zhou,X., Zhao,S., Dunn,D., von Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O., Weinstock,G. and Gibbs,R.A.

Direct Submission
Unpublished
2 (bases 1 to 285888)
Worley,K.C.
Direct Submission
Submitted (16-JAN-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 285888)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (22-SEP-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Sep 22, 2002 this sequence version replaced gi:21737099.
The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). As a result, the sequence may extend beyond the ends of the clone and there may be contigs that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GONW
Center clone name: CH230-38C17
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 239711 bases at least Q40
Consensus quality: 243503 bases at least Q30
Consensus quality: 245994 bases at least Q20
Estimated insert size: 277420; sum-of-contigs estimation
Quality coverage: 3x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)
* NOTE: This sequence may represent more than one clone.
* NOTE: This is a 'working draft' sequence. It currently consists of 6 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.
* 1 11520: contig of 11520 bp in length
* 11521 11620: gap of unknown length
* 11621 244182: contig of 232562 bp in length
* 244183 244282: gap of unknown length
* 244283 247660: contig of 3378 bp in length
* 247661 247761: gap of unknown length
* 247761 264056: contig of 16296 bp in length
* 264057 264156: gap of unknown length
* 264157 268805: contig of 4649 bp in length
* 268806 268905: gap of unknown length
* 268906 285888: contig of 16983 bp in length.
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Db 234124 CATGTAGTTGATGTTGATAGAG 234146

RESULT 5
BC070398/c
LOCUS
DEFINITION
Mus musculus peroxisome proliferator activator receptor delta, mRNA
(CDNA clone MGC:86084 IMAGE:5694282), complete cds.
ACCESSION
BC070398
VERSION
BC070398.1 GI:47125285
SOURCE
Mus musculus (house mouse)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
1 (bases 1 to 3191)
Strausberg,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G.,
Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
Altschul,S.F., Zeeberg,B., Buetow,K.H., Schaefer,C.P., Bhat,N.K.,
Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Hong,P.,
Diatchenko,K., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
Scapleton,M., Soares,M.B., Bonaldo,M.F., Casavant,T.L.,
Sheetz,T.E., Brownstein,M.J., Usdin,T.B., Toshiyuki,S.,
Carninci,P., Prange,C., Raha,S.S., Loquellano,N.A., Peters,G.J.,
Abramson,R.D., Mullaby,S.J., Bosak,S.A., McEwan,P.J.,
McKernan,K.J., Matek,A.A., Gunaratne,P.H., Richards,S.,
Worley,K.C., Hale,S., Garcia,A.M., Gay,L.J., Hulyk,S.W.,
Villalon,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A.,
Fahey,J., Helton,E., Kettman,M., Madan,A., Rodrigues,S.,
Sanchez,A., Whiting,M., Madan,A., Young,A.C., Shevchenko,Y.,
Bouffard,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D.,
Dickson,M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M.,
Schnurch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
Butterfield,Y.S., Krzywinski,M.I., Skalska,U., Smalish,D.E.,
Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences
Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)

PUBMED
12477932
2 (bases 1 to 3191)
Strausberg,R.
Direct Submission
Submitted (10-MAY-2004) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: cgapbs@mail.nih.gov
Tissue Procurement: Dr. Jim Lin, University of Iowa
cDNA Library Preparation: M. Bento Soares, University of Iowa
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: University of Iowa, Dr. M. Bento Soares and Dr.
Thomas L. Casavant.
Web site: http://genome.uiowa.edu
Contact: bento-soares@uiowa.edu; tom-casavant@uiowa.edu
Bonaldo,M.F., Akabogu,I., Bair,T., Bair,J., Crouch,K., Davis,A.,
Fiehler,K., Keppel,C., Kucaba,T., Lebeck,M., Melo,A., Schaefer,K.,
Scheetz,T., Smith,C., Snir,E., Tack,D., Trout,K., Walters,J.,
Casavant,T., Soares,M.B.
Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
Series: Plate: Row: Column: 0
This clone was selected for full length sequencing because it
passed the following selection criteria: Hexamer frequency ORF
analysis, Genomescan gene prediction.
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ORIGIN
Query Match      81.7%; Score 18.8; DB 10; Length 3191;
Best Local Similarity 90.9%; Pred. No. 3.1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CATGTATTTTCATGGGGATAGAG 22
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Db 1974 CATGTCTTTGAAGGGGATAGAG 1953

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RESULT 6
AB023160
LOCUS Homo sapiens mRNA for KIAA0943 protein, partial cds.
ACCESSION AB023160
VERSION AB023160.1 GI:4589529
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS Nagase, T., Ishikawa, K., Suyama, M., Kikuno, R., Hirose, M.,
Miyajima, N., Tanaka, A., Kotani, H., Nomura, N. and Ohara, O.
TITLE Prediction of the coding sequences of unidentified human genes.
XIII. The complete sequences of 100 new cDNA clones from brain
which code for large proteins in vitro
JOURNAL DNA Res. 6 (1), 63-70 (1999)
MEDLINE 99246063
PUBMED 10231032
REFERENCE 2 (bases 1 to 5634)
AUTHORS Ohara, O., Nagase, T. and Kikuno, R.
TITLE Direct Submission
JOURNAL Submitted (04-FEB-1999) Otsu DNA Research Institute,
Laboratory of DNA Technology; Yana 1532-3, Kiearazu, Chiba
292-0812, Japan (E-mail: cdnainfo@kazusa.or.jp, Tel: +81-438-52-3913,
Fax: +81-438-52-3914)
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Query Match 81.7%; Score 18.8; DB 9; Length 5634;
Best Local Similarity 90.9%; Pred. No. 3.1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 2 ATGTATTGATGGGATAGAGG 23
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Db 1556 ATATATTGATGGGAAAGAGG 1577
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RESULT 7
MMU420922/c
LOCUS Mus musculus partial PPARb/d gene for peroxisome proliferator
activated receptor beta/delta, exons 4-8.
DEFINITION
ACCESSION MMU420922
VERSION MMU420922.1 GI:22450050
KEYWORDS

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SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE 1
AUTHORS Larsen, L.K., Amri, E.Z., Mandrup, S., Pacot, C. and Kristiansen, K.
TITLE Genomic organization of the mouse peroxisome proliferator-activated
receptor beta/delta gene: alternative promoter usage and splicing
yield transcripts exhibiting differential translational efficiency
JOURNAL Biochem. J. 366 (Pt 3), 767-775 (2002)
MEDLINE 22201784
PUBMED 12059785
REFERENCE 2 (bases 1 to 8225)
AUTHORS Larsen, L.K.
TITLE Direct Submission
JOURNAL Submitted (12-SEP-2001) Larsen L.K., Basic, Center for clinical and
basic research, Ballerup Byvej 222, DK-2750 Ballerup, DENMARK
COMMENT Related sequences: AJ420918 - AJ420922.
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      Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 CATGTATTGATGGGGATAGAG 22
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DB      5116 CATGCTTTGAGGGGATAGAG 5095

RESULT 8
AC117508
LOCUS   AC117508
DEFINITION Homo sapiens 3 BAC RP11-207P12 (Roswell Park Cancer Institute Human
          BAC Library) complete sequence.
ACCESSION AC117508
VERSION   AC117508.5 GI:28882126
KEYWORDS  HTG.
SOURCE    Homo sapiens (human)
ORGANISM  Homo sapiens
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          Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D.,
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          Moore,S., Morgan,M., Moorish,T., Morris,S., Moser,M., Neal,D.,

```

Nelson, D., Newton, J., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokenkwo, S., Oguh, M., Okwuonu, G., Orsinger, N., Oviado, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Pickens, R., Primus, E., Pu, L.L., Quiles, M., Ren, Y., Rives, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Savery, G., Scherer, S., Scott, G., Shen, H., Shoochari, N., Sisson, I., Sodergren, E., Sonaite, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Svatek, A., Tabor, P., Tamarisa, A., Tamerisa, K., Tang, H., Tansey, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S., Usmani, K., Vaquez, L., Vera, V., Villalon, D., Vinson, R., Wang, Q., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S., Williams, G., Williamson, A., Wleczyk, R., Wooden, S., Worley, K., Wu, C., Wu, Y., Wu, Y. F., Zhou, J., Zorrilla, S., Naylor, S.L., Weinstein, G. and Gibbs, R.

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STSS are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL:

at ORD:
http://www.hgsc.bcm.tmc.edu:8088/quality.info/genbank.annotation.ht
ml.

FEATURES

source

Location/Qualifiers

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Query Match      81.7% Score 18.8; DB 9; Length 20838;
Best Local Similarity 90.9%; Pred. No. 3.2e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTTATTGATGGGATAGAGG 23
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Db 1078 ATGTTTTCATGGGATAGGG 1099

RESULT 9
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LOCUS      Homo sapiens 3 BAC RP11-722C17 (Roswell Park Cancer Institute Human
DEFINITION BAC Library) complete sequence.
ACCESSION  AC139617.6 GI:28557821
VERSION     HTG.
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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TITLE
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REFERENCE
AUTHORS
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JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Tabor,P., Tamerisa,A., Tamerisa,K., Tang,H., Tansey,J., Taylor,C.,
Taylor,T., Telford,B., Thomas,N., Thomas,S., Usmani,K., Vasquez,L.,
Vera,V., Villalón,D., Vinson,R., Wang,Q., Wang,S., Ward-Moore,S.,
Warren,R., Washington,C., Watlington,S., Williams,G.,
Williamson,A., Wleczyk,R., Wooden,S., Worley,K., Wu,C., Wu,Y.,
Wu,Y.F., Zhou,J., Zorrilla,S., Naylor,S.L., Weinstock,G. and
Gibbs,R.
Direct Submission
Unpublished
2 (bases 1 to 25901)
Worley,K.C.
Direct Submission
Submitted (07-FEB-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 25901)
Worley,K.C.
Direct Submission
Submitted (22-FEB-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 25901)
Worley,K.C.
Direct Submission
Submitted (25-FEB-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
INFORMATION: http://www.hgsc.bcm.tmc.edu/ or email
gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the
entire insert of this clone. Overlapping regions of clones are only
sequenced and submitted once, so the sequence for the remainder of
the insert may be found in the record for the adjacent clones.
Overlapping clones are noted at the beginning and end of the
Features listing.

ANNOTATION OF FEATURES:
STS are identified using ePCR (Genome Res. 7:541-550) searches
of a local database that includes entries from dbSTS, GDB, and
local mapping efforts.
Repeats are identified using RepeatMasker (A. Smit and P. Green,
unpublished.) for Human and Mouse sequences.
Genes and Region of sequence similarity are identified by BLAST
(Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the
EST and cDNA sequences. Genes demonstrate at least two exons
flanked by consensus splice sites that maintained sequence
continuity across the splice junctions. Sequences that are not
identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum
standard of double strand coverage with a minimum of 2 clones and 2
reads with no ambiguities or 2 chemistries with a minimum of 2
clones and 3 reads with no ambiguities. If the sequence quality for
a region does not meet this standard, it will be indicated in the
annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality
standards - estimated error rate less than 1 per 10,000 bases.
Reports of lowest quality individual bases and measures of base
quality are listed below. Description of the metrics can be found
at URL:
http://www.hgsc.bcm.tmc.edu:8088/quality.info/genbank.annotation.ht
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Query Match 81.7%; Score 18.8; DB 9; Length 25901;
Best Local Similarity 90.9%; Pred. No. 3.2e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTATTGATGGGATAGAGG 23
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Db 24889 ATGTTTTCATGGGATAGGG 24910
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RESULT 10
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LOCUS Homo sapiens chromosome 3 sequence from BAC 352H24 map 3q21 region
DEFINITION D3S3607-D3S1587, complete sequence.
ACCESSION AL449215
VERSION AL449215.3 GI:13752106
KEYWORDS HTG.
SOURCE Homo sapiens (human).
ORGANISM Homo sapiens
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Sudbrak,R., Ramser,J., Kosiura,A., Klueber,M., Borzym,K.,
Langer,I., Heitmann,K., Schuelzchen,S., Thompson,C., Lehrach,H. and
Reinhardt,R.
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 134296)
AUTHORS Direct Submission
TITLE Submitted (01-NOV-2000) MOLGENR, Abt. Lehrach, Max Planck Institut
JOURNAL Fuer Molekulare Genetik, Innestrasse 73, Berlin, 14195 Germany
COMMENT On Apr 22, 2001 this sequence version replaced gi:12311590.
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db_xref="taxon:9606"
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library"
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ORIGIN
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Best Local Similarity 90.9%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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RESULT 11
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LOCUS Homo sapiens BAC clone RP11-367H1 from 2, complete sequence.
DEFINITION
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ACCESSION      AC133528
VERSION        AC133528.4
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SOURCE         Homo sapiens (human)
ORGANISM       Homo sapiens
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               Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 151700)
AUTHORS        Sulston,J.E. and Waterston,R.
TITLE          Toward a complete human genome sequence
JOURNAL        Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE        99063792
PUBMED         9847074
REFERENCE      2 (bases 1 to 151700)
AUTHORS        Ali,J., Haakenson,W., Dignan,G. and Doebber,A.
TITLE          The sequence of Homo sapiens BAC clone RP11-367H1
JOURNAL        Unpublished (2001)
REFERENCE      3 (bases 1 to 151700)
AUTHORS        Waterston,R.H.
TITLE          Direct Submission
JOURNAL        Submitted (13-SEP-2002) Genome Sequencing Center, Washington
               University School of Medicine, 4444 Forest Park Parkway, St. Louis,
               MO 63108, USA
REFERENCE      4 (bases 1 to 151700)
AUTHORS        Waterston,R.H.
TITLE          Direct Submission
JOURNAL        Submitted (23-OCT-2002) Genome Sequencing Center, Washington
               University School of Medicine, 4444 Forest Park Parkway, St. Louis,
               MO 63108, USA
REFERENCE      5 (bases 1 to 151700)
AUTHORS        Waterston,R.
TITLE          Direct Submission
JOURNAL        Submitted (27-NOV-2002) Department of Genetics, Washington
               University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
REFERENCE      6 (bases 1 to 151700)
AUTHORS        Waterston,R.H.
TITLE          Direct Submission
JOURNAL        Submitted (18-JAN-2003) Genome Sequencing Center, Washington
               University School of Medicine, 4444 Forest Park Parkway, St. Louis,
               MO 63108, USA
REFERENCE      7 (bases 1 to 151700)
AUTHORS        Waterston,R.
TITLE          Direct Submission
JOURNAL        Submitted (19-FEB-2003) Department of Genetics, Washington
               University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
COMMENT        On Jan 18, 2003 this sequence version replaced gi:25777548.
               ----- Genome Center
               Center: Washington University Genome Sequencing Center
               Center code: WUGSC
               Web site: http://genome.wustl.edu/gsc
               Contact: sapiens@wustl.wustl.edu
               ----- Summary Statistics
               -----
               Center project name: H_NH0367H01

```

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osoegawa,K., Woon,P.Y., Zhao,B., Frengen,E., Tatenio,M., Catanese,J.J. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at <http://www.chori.org>
VECTOR: pBACE3.6

NEIGHBORING SEQUENCE INFORMATION:

This sequence is not the entire insert of the clone. This clone is overlapped by AC110299.

Discrepant bases between AC114730, AC132150 and clone sequence.

Data from AC132150 and AC114730 was used to finish this clone.

FEATURES

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1656..1840	repeat_region
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repeat_region /rpt_family="(CA)n"
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repeat_region /rpt_family="L1"
repeat_region 13548. .13781
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repeat_region 13782. .13843
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repeat_region 13844. .14141
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Best Local Similarity 90.9%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 ATGATTGATGGGATAGG 23
Db 42701 ATATTGATGGGAAAGG 42680

RESULT 12
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LOCUS AC102466 163947 bp DNA linear HTG 16-SEP-2004
DEFINITION Mus musculus chromosome 3 clone RP24-337A2 map 3, *** SEQUENCING IN
PROGRESS ***, 5 unordered pieces.
ACCESSION AC102466
VERSION AC102466.6 GI:52138860
KEYWORDS HTG; HTGS PHASE1; HTGS FULLTOP; HTGS ACTIVEFIN.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

```

REFERENCE

1 (bases 1 to 163947)
 Birren, B., Nusbaum, C., and Lander, E.
 Mus musculus chromosome 3, clone RP24-337A2
 unpublished
 2 (bases 1 to 163947)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,
 Anderson, S., Barna, N., Bastien, V., Boguslavskiy, L., Boukhgalter, B.,
 Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B.,
 Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A.,
 Cooke, P., DeArellano, K., Dewar, K., Diaz, J.S., Dodge, S., Faro, S.,
 Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Grand-Pierre, N.,
 Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,
 Hagos, B., Heaford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R.,
 Jones, C., Kamat, A., Karatas, A., Kells, C., LaRocque, K.,
 Lamazares, R., Landers, T., Lechoczy, J., Levine, R., Liu, G.,
 MacLean, C., MacDonald, P., Major, J., Marquis, N., Matthews, C.,
 McCarthy, M., McEwan, P., McKernan, K., McPheeters, R., Meldrum, J.,
 Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C.,
 Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D.,
 Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V.,
 Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P.,
 Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schuback, R.,
 Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
 Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,
 Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H.,
 Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G.,
 Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission
 Submitted (23-NOV-2001) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 3 (bases 1 to 163947)

Birren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N.,
 Anderson, M., Anderson, S., Atacchi, H.M., Barna, N., Bastien, V.,
 Bloom, T., Boguslavskiy, L., Boukhgalter, B., Camarata, J., Chang, J.,
 Choepel, Y., Collymore, A., Cook, A., Cooke, P., Corum, B.,
 DeArellano, K., Diaz, J.S., Dodge, S., Dooley, K., Dorris, L.,
 Erickson, J., Faro, S., Ferreira, P., FitzGerald, M., Gage, D.,
 Galagan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N.,
 Hagojian, D., Hagos, B., Hall, J., Horton, L., Hulme, W., Iliev, I.,
 Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T.,
 Levine, R., Lindblad-Toh, K., Liu, G., Liu, X., Lui, A., Mabbitt, R.,
 MacLean, C., MacDonald, P., Major, J., Manning, J., Matthews, C.,
 McCarthy, M., Meldrum, J., Meneus, L., Mihova, T., Mlenga, V.,
 Murphy, T., Naylor, J., Nguyen, C., Nguyen, T., Nicol, R., Norbu, C.,
 O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
 Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C.,
 Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schuback, R.,
 Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N.,
 Stojanovic, N., Stubbs, M., Talamas, J., Tesfaye, S., Theodore, J.,
 Topham, K., Travers, M., Vassiliev, H., Venkataraman, V.S., Viel, R.,
 Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L.,
 Zimmer, A. and Zody, M.

Direct Submission
 Submitted (16-SEP-2004) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Sep 16, 2004 this sequence version replaced gi:51536822.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html
 ----- Genome Center
 Center: Whitehead Institute/MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@broad.mit.edu
 ----- Project Information
 Center project name: L18861
 Center clone name: 337_A_2

* NOTE: This is a 'working draft' sequence. It currently
 * consists of 5 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as

REFERENCE

1 (bases 1 to 163947)
 Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N.,
 Anderson, S., Barna, N., Bastien, V., Boguslavskiy, L., Boukhgalter, B.,
 Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B.,
 Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A.,
 Cooke, P., DeArellano, K., Dewar, K., Diaz, J.S., Dodge, S., Faro, S.,
 Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Grand-Pierre, N.,
 Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,
 Hagos, B., Heaford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R.,
 Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T.,
 Levine, R., Lindblad-Toh, K., Liu, G., Liu, X., Lui, A., Mabbitt, R.,
 MacLean, C., MacDonald, P., Major, J., Manning, J., Matthews, C.,
 McCarthy, M., Meldrum, J., Meneus, L., Mihova, T., Mlenga, V.,
 Murphy, T., Naylor, J., Nguyen, C., Nguyen, T., Nicol, R., Norbu, C.,
 O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
 Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C.,
 Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schuback, R.,
 Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N.,
 Stojanovic, N., Stubbs, M., Talamas, J., Tesfaye, S., Theodore, J.,
 Topham, K., Travers, M., Vassiliev, H., Venkataraman, V.S., Viel, R.,
 Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L.,
 Zimmer, A. and Zody, M.

Direct Submission
 Submitted (16-SEP-2004) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Sep 16, 2004 this sequence version replaced gi:51536822.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html
 ----- Genome Center
 Center: Whitehead Institute/MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@broad.mit.edu
 ----- Project Information
 Center project name: L18861
 Center clone name: 337_A_2

* NOTE: This is a 'working draft' sequence. It currently
 * consists of 5 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

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 * 70702 96871: contig of 26170 bp in length
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FEATURES

source

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 /db_xref="taxon:10090"
 /chromosome="3"
 /map="3"
 /clone="RP24-337A2"
 /clone_lib="RPCI-24 Male Mouse BAC"

ORIGIN

Query Match 81.7%; Score 18.8; DB 2; Length 163947;
 Best Local Similarity 90.9%; Pred. No. 3.4e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTATTGATGGGATAGAGG 23

Db 42059 ATTATTGATGGGAGAGGG 42080

RESULT 13

AC151841

LOCUS AC151841 167797 bp DNA linear HTG 06-OCT-2004
 DEFINITION Mus musculus chromosome 7 clone RP23-128P10, *** SEQUENCING IN
 PROGRESS ***, 63 unordered pieces.

AC151841

VERSION AC151841.1 GI:53828886

KEYWORDS HTG; HTGS PHASE1.

SOURCE Mus musculus (house mouse)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 Wilson, R.K.
 1 (bases 1 to 167797)

REFERENCE

AUTHORS The sequence of Mus musculus clone

TITLE Unpublished

JOURNAL 2 (bases 1 to 167797)

REFERENCE Wilson, R.K.

AUTHORS Direct Submission

TITLE Submitted (06-OCT-2004) Genome Sequencing Center, 4444 Forest Park

JOURNAL Parkway, St. Louis, MO 63108, USA

COMMENT

----- Genome Center -----
 Center: Washington University Genome Sequencing Center
 Center code: WUGSC
 Web site: http://genome.wustl.edu
 ----- Project Information -----
 Center project name: M_BA0128P10
 ----- Summary Statistics -----
 Sequencing vector: M13; 0%

Chemistry: Dye-terminator Big Dye; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.990319

Consensus quality: 132837 bases at least Q40
 Consensus quality: 141875 bases at least Q30
 Consensus quality: 146026 bases at least Q20

 * NOTE: This is a 'working draft' sequence. It currently

* consists of 63 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

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 * 4105 4204: gap of unknown length
 * 4205 5597: contig of 1393 bp in length
 * 5598 5697: gap of unknown length
 * 5698 7028: contig of 1231 bp in length
 * 7029 8365: contig of 1337 bp in length
 * 8366 8465: gap of unknown length
 * 8466 9878: contig of 1413 bp in length
 * 9879 9979 11628: contig of 1650 bp in length
 * 11629 11728: gap of unknown length
 * 11729 13119: contig of 1391 bp in length
 * 13120 13219: gap of unknown length
 * 13220 14722: contig of 1503 bp in length
 * 14723 14822: gap of unknown length
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 * 19280 20692: contig of 1413 bp in length
 * 20693 20792: gap of unknown length
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 * 25284 26438: contig of 1155 bp in length
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 * 37275 38990: contig of 1716 bp in length
 * 38991 39090: gap of unknown length
 * 39091 40783: contig of 1693 bp in length
 * 40784 40883: gap of unknown length
 * 40884 42469: contig of 1586 bp in length
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 * 42570 44071: contig of 1502 bp in length
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* 100140 105686: contig of 5546 bp in length
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* 105787 111728: contig of 5942 bp in length
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* 111828 116854: contig of 5026 bp in length
* 116854 116954: gap of unknown length
* 116954 123034: contig of 6080 bp in length
* 123034 123134: gap of unknown length
* 123134 128962: contig of 5828 bp in length
* 128962 129062: gap of unknown length
* 129062 133910: contig of 4848 bp in length
* 133910 134011: contig of 5078 bp in length
* 134011 139089: gap of unknown length
* 139089 145581: contig of 6393 bp in length
* 145581 145681: gap of unknown length
* 145681 153130: contig of 7449 bp in length
* 153130 153230: gap of unknown length
* 153230 153350: contig of 6120 bp in length
* 153350 159351: gap of unknown length
* 159351 167797: contig of 8347 bp in length.
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misc_feature 1411. .2848
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16206. .17617

Query Match 81.7%; Score 18.8; DB 2; Length 167797;
Best Local Similarity 90.9%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTAATTTGATGGGATAGAGG 23
Db 43214 ATGTAATTTGATGGGATAGAGG 43235

RESULT 14
AC118744
LOCUS AC118744
DEFINITION Mus musculus chromosome 6 clone RP24-182M11 map 6, *** SEQUENCING
IN PROGRESS ***, 7 unordered pieces.
ACCESSION AC118744
VERSION AC118744.5 GI:42761784
KEYWORDS HTG: HTGS_PHASE1; HTGS_FULLTOP; HTGS_ACTIVEPIN.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 171446)
AUTHORS Birren,B., Nusbaum,C. and Lander,E.
JOURNAL Unpublished
TITLE Mus musculus chromosome 6, clone RP24-182M11
REFERENCE 2 (bases 1 to 171446)
AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L.,
Boukhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J.,
Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
Kamat,A., Karatas,A., Kells,C., LaRocque,K., Lamazares,R.,
Landers,T., Lechoczky,J., Levine,R., Lindblad-Toh,K., Liu,G.,
Maclean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C.,
McCarthy,M., McEwan,P., McKernan,K., Meldrim,J., Meneus,L.,
Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R.,
Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D.,
Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupack,R.,
Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
TITLE Direct Submission
JOURNAL Submitted (20-APR-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE 3 (bases 1 to 171446)
AUTHORS Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N.,
Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T.,
Boguslavsky,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y.,
Collymore,A., Cook,A., Cooke,P., Corum,B., DeArellano,K.,

```


Boguslavskiy, L., Boukhgalter, B., Camarata, J., Chang, J., Choepel, Y., Collymore, A., Cook, A., Cooke, P., Corum, B., DeArellano, K., Diaz, J. S., Dodge, S., Dooley, K., Dorris, L., Erickson, J., Faro, S., Ferreira, P., FitzGerald, M., Gage, D., Galagan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N., Hagopian, D., Hagos, B., Hall, J., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, G., Lui, X., Mabbitt, R., Matthews, C., McDonald, P., Major, J., Manning, J., Menus, L., Mhova, T., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schupback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Stubbs, M., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Venkataraman, V. S., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission
Submitted (30-SEP-2003) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
6 (bases 1 to 175280)

TITLE

JOURNAL

REFERENCE

AUTHORS

Barren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N., Anderson, M., Arachchi, H. M., Barna, N., Bastien, V., Bloom, T., Boguslavskiy, L., Boukhgalter, B., Camarata, J., Chang, J., Choepel, Y., Collymore, A., Cook, A., Cooke, P., Corum, B., DeArellano, K., Diaz, J. S., Dodge, S., Dooley, K., Dorris, L., Erickson, J., Faro, S., Ferreira, P., FitzGerald, M., Gage, D., Galagan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N., Hagopian, D., Hagos, B., Hall, J., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, G., Lui, X., Mabbitt, R., Matthews, C., McDonald, P., Major, J., Manning, J., Menus, L., Mhova, T., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schupback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Stubbs, M., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Venkataraman, V. S., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission
Submitted (24-OCT-2003) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Sep 10, 2003 this sequence version replaced gi:33147222.
All repeats were identified using RepeatMasker:
Smit, A. F. A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

TITLE

JOURNAL

COMMENT

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L22700
Center clone name: 375_O_10

FEATURES

source

Query Match 81.7%; Score 18.8; DB 10; Length 175280;
Best Local Similarity 90.9%; Pred. No. 3.4e+02; Indels 0; Gaps 0;
Matches 20; Conservative 0; Mismatches 2;

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12868. .12942

Query Match

Best Local Similarity

Matches 20; Conservative 0; Mismatches 2;

QY

DB

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Job time : 740.234 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 12, 2005, 22:21:57 ; Search time 184.245 Seconds
(without alignments)
738.985 Million cell updates/sec

Title: US-10-673-854-1

Perfect score: 23

Sequence: 1 catgtatttgatggggatagagg 23

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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4: Geneseqn2001as:*
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13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	18.8	81.7	5651	4 ABA08791	Aba08791 Human dJ8
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4	18.8	81.7	5654	4 AAh99496	Aah99496 Human pro
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11	18.2	79.1	1917	6 ABV75719	Abv75719 Human B-C
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13	18.2	79.1	29111	11 ACN44424	Acn44424 Mouse gen
14	18	78.3	2286	13 ADS46987	Ads46987 Bacterial
15	17.8	77.4	238	5 ABV12937	Abv12937 Human pro
16	17.8	77.4	238	5 ABV03768	Abv03768 Human pro
17	17.8	77.4	416	5 ABV34062	Abv34062 Human pro
18	17.8	77.4	745	4 AAI93090	Aai93090 Human pol
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	23	17.8	77.4	5975	6	ABL32236	Ab132236 Human imm
	24	17.8	77.4	11046	6	ABK31536	Abk31536 Signal tr
	25	17.2	74.8	33	6	ABA95678	Aba95678 Human int
	26	17.2	74.8	648	10	ADD33537	Add33537 Mouse mit
c	27	17.2	74.8	957	6	ABN98549	Abn98549 Arabidops
	28	17.2	74.8	1026	4	AAH52942	Aah52942 S. epider
	29	17.2	74.8	2250	4	AAH52372	Aah52372 S. epider
	30	17.2	74.8	2430	8	ACA46750	Ac46750 Prokaryot
	31	17.2	74.8	2448	6	ABN91314	Abn91314 Staphyloc
	32	17.2	74.8	2448	13	ADS01065	Ad01065 Staphyloc
	33	17.2	74.8	3427	4	AAH54925	Aah54925 S. epider
c	34	17.2	74.8	4198	4	AAH54240	Aah54240 S. epider
	35	17.2	74.8	8700	6	ABL33003	Ab133003 Human imm
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	38	17.2	74.8	110000	12	ADQ97331	Adq97331 2
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	45	16.8	73.0	5757	4	ABL08602	Ab108602 Drosophil

ALIGNMENTS

RESULT 1

ABA08791

ID ABA08791 standard; cDNA; 5651 BP.

AC ABA08791;

XX 11-JAN-2002 (first entry)

XX Human dJ889M15.3 homologue-encoding cDNA, SEQ ID NO:567.

XX Human; cytokine; cell proliferation; cell differentiation; growth factor;
KW haematopoiesis regulation; tissue growth; immunomodulator; activin;
KW inhibitor; chemotaxis; chemokinesis; thrombolysis; oncogenesis;
KW proliferation; metastasis; cancer; tumour; haematopoietic disorder;
KW myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;
KW chronic inflammatory condition; proliferative retinopathy;
KW atherosclerosis; coronary heart disease; arterial ischaemia;
KW bone disorder; osteoporosis; vascular growth disorder;
KW tissue regeneration; wound healing; infection; immune disorder;
KW cell culture; drug screening; gene therapy; antiinflammatory;
KW antiasthmatic; antiarthritic; haemostatic; antiarteriosclerotic;
KW cytotatic; osteopathic; vasotropic; cardiant; virucide; antibacterial;
KW antifungal; vulnery; antitumor; ss.

XX Homo sapiens.

XX WO200157188-A2.

PD 09-AUG-2001.

XX 05-FEB-2001; 2001WO-US003800.

PF 03-FEB-2000; 2000US-00496914.

PR 27-APR-2000; 2000US-00560875.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT;

XX WPI; 2001-457740/49.

XX P-PSDB; ABB11547.

PT Human proteins and DNA encoding sequences useful for preventing, treating

PT or ameliorating a medical condition in a mammalian subject e.g. arthritis
 XX and cancer.

PS Claim 1; Page 582-584; 1963pp; English.

XX Sequences ABB10981-ABB12330 represent 1350 novel human polypeptides, and
 CC sequences ABA08225-ABA09574 represent nucleic acids encoding them. The
 CC invention also relates to vectors and recombinant host cells comprising a
 CC nucleotide of the invention, methods of producing the novel polypeptides,
 CC antibodies against the polypeptides, methods of detecting the nucleotides
 CC or polypeptides in a sample, and methods of identifying compounds which
 CC bind to polypeptides of the invention. Although novel, many of the
 CC polypeptides of the invention have homology to known proteins, thereby
 CC giving an insight into their probable biological activities, and hence
 CC potential therapeutic applications. The polypeptides of the invention may
 CC have various activities, including cytokine, cell proliferation or cell
 CC differentiation activities; stem cell growth factor activity;
 CC haematopoiesis regulatory activity; tissue growth activity;
 CC immunomodulatory activity; activin- or inhibin-related activities;
 CC chemotactic or chemokinetic activities; haemostatic, thrombotic or
 CC thrombolytic activities; receptor or ligand activities; or may be
 CC involved in oncogenesis, cancer cell proliferation or metastasis.
 CC Depending on their biological activities, polypeptides and nucleotides of
 CC the invention are useful for preventing, treating or ameliorating medical
 CC conditions, e.g., by protein or gene therapy. Such conditions include
 CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell
 CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),
 CC proliferative retinopathy, atherosclerosis, coronary heart disease,
 CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal
 CC vascular growth. Polypeptides involved with tissue regeneration and
 CC repair (or nucleic acids encoding them) may be used to promote wound
 CC healing (e.g., of burns, incisions and ulcers), while those with
 CC immunomodulatory activities may be used in the treatment of viral,
 CC bacterial and fungal infections in addition to immune disorders.
 CC Polypeptides with growth factor activity may be used in cell cultures to
 CC promote cell growth. For example, such polypeptides may be used to
 CC manipulate stem cells in culture to give rise to neuroepithelial cells
 CC that can be used to augment or replace cells damaged by illness,
 CC autoimmune disease or accidental damage. The polypeptides and nucleotides
 CC may also be used in the diagnosis of the above conditions, and in drug
 CC screening techniques. The present sequence represents a cDNA encoding a
 CC novel human polypeptide of the invention

XX Sequence 5651 BP; 1077 A; 1633 C; 1680 G; 1261 T; 0 U; 0 Other;

Query Match 81.7%; Score 18.8; DB 4; Length 5651;
 Best Local Similarity 90.9%; Pred. No. 1e-02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTATTTCATGGGATAGAGG 23

Db 1556 ATATATTTCATGGGAAAGAGG 1577

RESULT 2

AAH99780/c

ID AAH99780 standard; cDNA; 5653 BP.

XX AAH99780;

DT 16-OCT-2001 (first entry)

DE Human protein encoding cDNA sequence SEQ ID NO:615.

XX Human; cancer; ulcer; HIV infection; human immunodeficiency virus;
 KW antiinflammatory; antirheumatic; antiarthritic; immunosuppressive;
 KW antibacterial; endocrine; cardiant; central nervous system; virucide;
 KW anti-HIV; fungicide; antitumor; cardiovascular; antianaemic; anaemia;
 KW antiaggregant; haemostatic; vulnerary; antileuk; osteopathic; eczema;
 KW dermatological; antiallergic; antiasthmatic; antidiabetic; cytostatic;
 KW neuroprotective; nootropic; antiparkinsonian; infection;
 KW immunostimulant; gene therapy; antisense therapy; vaccine; inflammation;
 KW antianaphylactic; rheumatoid arthritis; septic shock; pancreatitis;

KW cardiac dysfunction; neuropathology; cardiac anaphylaxis; autoimmunity;
 KW genetic disease; haematopoietic disorder; platelet disorder; asthma;
 KW thrombocytopaenia; osteoporosis; severe combined immunodeficiency;
 KW allergic rhinitis; diabetes; multiple sclerosis; depression;
 KW Alzheimer's disease; Parkinson's disease; neurodegenerative disorder;
 KW neurological disorder; ps.

XX Homo sapiens.

PN WO200153455-A2.

XX 26-JUL-2001.

XX 22-DEC-2000; 2000WO-US035017.

XX 23-DEC-1999; 99US-00471275.

PR 21-JAN-2000; 2000US-00488725.

PR 25-APR-2000; 2000US-00552317.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT;

XX WPI; 2001-457603/49.

XX P-PSDB; AAM25839.

XX Isolated human polynucleotides encoding polypeptides, useful for the
 PT treatment and diagnosis of e.g. cancer, ulcers and HIV infection.

XX Claim 1; Page 642-644; 1217pp; English.

XX AAH99166 to AAH99904 encode the human proteins given in AAM25225 to
 CC AAM25963. The proteins can have activities based on the tissues and cells
 CC they are expressed in, such as: antiinflammatory; antirheumatic;
 CC antiarthritic; immunosuppressive; antibacterial; endocrine; cardiant;
 CC central nervous system; virucide; anti-HIV; fungicide; antitumor;
 CC cardiovascular; antianaemic; antiaggregant; haemostatic; vulnerary;
 CC antileuk; osteopathic; dermatological; antiallergic; antiasthmatic;
 CC antidiabetic; cytostatic; neuroprotective; antidepressant; nootropic;
 CC antiparkinsonian; and immunostimulant. The proteins and polynucleotides
 CC encoding them can be used in gene therapy, antisense therapy and vaccine
 CC production. The proteins and polynucleotides are useful for screening for
 CC agonists or antagonists of a protein and for the treatment and diagnosis
 CC of disorders associated with the activity of a protein e.g. inflammation,
 CC rheumatoid arthritis, septic shock, pancreatitis, cardiac dysfunction,
 CC neuropathology, cardiac anaphylaxis, viral, bacterial, HIV and fungal
 CC infections, autoimmunity, genetic diseases, haematopoietic disorders,
 CC anaemia, platelet disorders, thrombocytopaenia, wounds, ulcers, eczema,
 CC osteoporosis, severe combined immunodeficiency, eczema, allergic
 CC rhinitis, asthma, diabetes, cancer, multiple sclerosis, depression,
 CC Alzheimer's disease, Parkinson's disease, neurodegenerative and
 CC neurological disorders

XX Sequence 5653 BP; 1262 A; 1680 C; 1632 G; 1079 T; 0 U; 0 Other;

Query Match 81.7%; Score 18.8; DB 4; Length 5653;
 Best Local Similarity 90.9%; Pred. No. 1e-02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTATTTCATGGGATAGAGG 23

Db 4098 ATATATTTCATGGGAAAGAGG 4077

RESULT 3

AAI60854/c

ID AAI60854 standard; cDNA; 5653 BP.

XX AAI60854;

XX 22-OCT-2001 (first entry)

DE Human polynucleotide SEQ ID NO 4843.

XX Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer;
 KW peripheral nervous system; neuropathy; central nervous system; CNS;
 KW Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
 KW amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
 KW chemokine; thrombolytic; drug screening; arthritis; inflammation;
 KW leukaemia; ss.
 XX Homo sapiens.
 XX WO200153312-A1.
 XX 26-JUL-2001.
 XX 26-DEC-2000; 2000WO-US034263.
 XX 23-DEC-1999; 99US-00471275.
 PR 21-JAN-2000; 2000US-0048725.
 PR 25-APR-2000; 2000US-00552317.
 PR 20-JUN-2000; 2000US-00598042.
 PR 19-JUL-2000; 2000US-00620312.
 PR 03-AUG-2000; 2000US-00653450.
 PR 14-SEP-2000; 2000US-00662191.
 PR 19-OCT-2000; 2000US-00693036.
 PR 29-NOV-2000; 2000US-00727344.
 XX (HYSE-) HYSEQ INC.
 XX Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;
 PI Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J, Zhao QH;
 PI Zhou P, Goodrich R, Drmanac RT;
 XX WPI; 2001-442253/47.
 DR P-PSDB; AAM41698.
 XX Novel nucleic acids and polypeptides, useful for treating disorders such
 PT as central nervous system injuries.
 XX Claim 1; SEQ ID NO 4843; 10078pp; English.
 XX The invention relates to human nucleic acids (AA157798-AA161369) and the
 CC encoded polypeptides (AAM38642-AAM42213) with nootropic,
 CC immunosuppressant and cytostatic activity. The polynucleotides are useful
 CC in gene therapy. A composition containing a polypeptide or polynucleotide
 CC of the invention may be used to treat diseases of the peripheral nervous
 CC system, such as peripheral nervous injuries, peripheral neuropathy and
 CC localised neuropathies and central nervous system diseases, such as
 CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
 CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
 CC utilisation of the activities such as: Immune system suppression,
 CC Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic
 CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
 CC assays for receptor activity, arthritis and inflammation, leukaemias and
 CC C.N.S disorders. Note: The sequence data for this patent did not form
 CC part of the printed specification
 XX Sequence 5653 BP; 1262 A; 1680 C; 1632 G; 1079 T; 0 U; 0 Other;
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 Query Match 81.7%; Score 18.8; DB 4; Length 5653;
 Best Local Similarity 90.9%; Pred. No. 1e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 2 ATGTATTGATGGGATAGG 23
 Db 4098 ATATATTGATGGGAAGAGG 4077
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 ID AAH99496 standard; cDNA; 5654 BP.
 XX
 AC AAH99496;
 XX

DT 16-OCT-2001 (first entry)
 XX Human protein encoding cDNA sequence SEQ ID NO:331.
 DE Human; cancer; ulcer; HIV infection; human immunodeficiency virus;
 XX antinflammatory; antirheumatic; antiarthritic; immunosuppressive;
 KW antibacterial; endocrine; cardiant; central nervous system; virucide;
 KW anti-HIV; fungicide; antimutagen; cardiovascular; anaemia;
 KW antiaggregant; haemostatic; vulnary; antilucer; osteopathic; eczema;
 KW dermatological; antiallergic; antiasthmatic; antidiabetic; cytostatic;
 KW neuroprotective; antidepressant; nootropic; antiparkinsonian; infection;
 KW immunostimulant; gene therapy; antisense therapy; vaccine; inflammation;
 KW antianaphylactic; rheumatoid arthritis; septic shock; pancreatitis;
 KW cardiac dysfunction; neuropathology; cardiac anaphylaxis; autoimmunity;
 KW genetic disease; haematopoietic disorder; platelet disorder; asthma;
 KW thrombocytopaenia; osteoporosis; severe combined immunodeficiency;
 KW allergic rhinitis; diabetes; multiple sclerosis; depression; disorder;
 KW Alzheimer's disease; Parkinson's disease; neurodegenerative disorder;
 KW neurological disorder; ss.
 OS Homo sapiens.
 XX WO200153455-A2.
 XX 26-JUL-2001.
 XX 22-DEC-2000; 2000WO-US035017.
 XX 23-DEC-1999; 99US-00471275.
 PR 21-JAN-2000; 2000US-00488725.
 PR 25-APR-2000; 2000US-00552317.
 XX (HYSE-) HYSEQ INC.
 XX Tang YT, Liu C, Drmanac RT;
 PI WPI; 2001-457603/49.
 DR P-PSDB; AAM25555.
 XX Isolated human polynucleotides encoding polypeptides, useful for the
 PT treatment and diagnosis of e.g. cancer, ulcers and HIV infection.
 PT Claim 1; Page 444-446; 1217pp; English.
 PS AAH99166 to AAH99904 encode the human proteins given in AAM25225 to
 CC AAM25963. The proteins can have activities based on the tissues and cells
 CC they are expressed in, such as: antiinflammatory; antirheumatic;
 CC antarthritic; immunosuppressive; antibacterial; endocrine; cardiant;
 CC central nervous system; virucide; anti-HIV; fungicide; antimutagen;
 CC cardiovascular; anaemia; antiaagregant; haemostatic; vulnary;
 CC antilucer; osteopathic; dermatological; antiallergic; antiasthmatic;
 CC antidiabetic; cytostatic; neuroprotective; antidepressant; nootropic;
 CC antiparkinsonian; and immunostimulant. The proteins and polynucleotides
 CC encoding them can be used in gene therapy, antisense therapy and vaccine
 CC production. The proteins and polynucleotides are useful for screening for
 CC agonists or antagonists of a protein and for the treatment and diagnosis
 CC of disorders associated with the activity of a protein e.g. inflammation,
 CC rheumatoid arthritis, septic shock, pancreatitis, cardiac dysfunction,
 CC neuropathology, cardiac anaphylaxis, viral, bacterial, HIV and fungal
 CC infections, autoimmunity, genetic diseases, haematopoietic disorders,
 CC anaemia, platelet disorders, thrombocytopaenia, wounds, burns, ulcers,
 CC osteoporosis, severe combined immunodeficiency, eczema, allergic
 CC rhinitis, asthma, diabetes, cancer, multiple sclerosis, depression,
 CC Alzheimer's disease, Parkinson's disease, neurodegenerative and
 CC neurological disorders
 XX Sequence 5654 BP; 1078 A; 1631 C; 1682 G; 1263 T; 0 U; 0 Other;
 SQ
 Query Match 81.7%; Score 18.8; DB 4; Length 5654;
 Best Local Similarity 90.9%; Pred. No. 1e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 2 ATGTATTGATGGGATAGG 23

Db 1559 ATATATTGATGGGGAAGG 1580
|||||
RESULT 5
ABA08796
ID ABA08796 standard; cDNA; 5665 BP.
XX ABA08796;
XX
DT 11-JAN-2002 (first entry)
XX
DE Human dJ889M15.3 homologue-encoding cDNA, SEQ ID NO:572.
XX
KW Human; cytokines; cell proliferation; cell differentiation; growth factor;
KW haematopoiesis regulation; tissue growth; immunomodulator; activin;
KW inhibin; chemotaxis; chemokinesis; thrombolysis; oncogenesis;
KW proliferation; metastasis; cancer; tumour; haematopoietic disorder;
KW myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;
KW chronic inflammatory condition; proliferative retinopathy;
KW atherosclerosis; coronary heart disease; arterial ischaemia;
KW bone disorder; osteoporosis; vascular growth disorder;
KW tissue regeneration; wound healing; infection; immune disorder;
KW cell culture; drug screening; gene therapy; antiinflammatory;
KW antiasthmatic; antiarthritic; haemostatic; antiarteriosclerotic;
KW cytostatic; osteopathic; vasotropic; cardiant; virucide; antibacterial;
KW antifungal; vulnery; antiulcer; ss.
XX
OS Homo sapiens.
XX
PN WO200157188-A2.
XX
PD 09-AUG-2001.
XX
PF 05-FEB-2001; 2001WO-US003800.
XX
PR 03-FEB-2000; 2000US-00496914.
PR 27-APR-2000; 2000US-00560875.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Tang YT, Liu C, Drmanac RT;
XX
XX WPI; 2001-457740/49.
DR P-PSDB; ABB11552.
XX
PT Human proteins and DNA encoding sequences useful for preventing, treating
PT or ameliorating a medical condition in a mammalian subject e.g. arthritis
PT and cancer.
XX
PS Claim 1; Page 586-587; 1963pp; English.
XX
CC Sequences ABB10981-ABB12330 represent 1350 novel human polypeptides, and
CC sequences ABA08225-ABA09574 represent nucleic acids encoding them. The
CC invention also relates to vectors and recombinant host cells comprising a
CC nucleotide of the invention, methods of producing the novel polypeptides,
CC antibodies against the polypeptides, methods of detecting the nucleotides
CC or polypeptides in a sample, and methods of identifying compounds which
CC bind to polypeptides of the invention. Although novel, many of the
CC polypeptides of the invention have homology to known proteins, thereby
CC giving an insight into their probable biological activities, and hence
CC potential therapeutic applications. The polypeptides of the invention may
CC have various activities, including cytokine, cell proliferation or cell
CC differentiation activities; stem cell growth factor activity;
CC haematopoiesis regulatory activity; tissue growth activity;
CC immunomodulatory activity; activin- or inhibin-related activities;
CC chemotactic or chemokinetic activities; haemostatic, thrombotic or
CC thrombolytic activities; receptor or ligand activities; or may be
CC involved in oncogenesis, cancer cell proliferation or metastasis.
CC Depending on their biological activities, polypeptides and nucleotides of
CC the invention are useful for preventing, treating or ameliorating medical
CC conditions, e.g., by protein or gene therapy. Such conditions include
CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell

CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),
CC proliferative retinopathy, atherosclerosis, coronary heart disease,
CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal
CC vascular growth. Polypeptides involved with tissue regeneration and
CC repair (or nucleic acids encoding them) may be used to promote wound
CC healing (e.g., of burns, incisions and ulcers), while those with
CC immunomodulatory activities may be used in the treatment of viral,
CC bacterial and fungal infections in addition to immune disorders.
CC Polypeptides with growth factor activity may be used in cell cultures to
CC promote cell growth. For example, such polypeptides may be used to
CC manipulate stem cells in culture to give rise to neuroepithelial cells
CC that can be used to augment or replace cells damaged by illness.
CC autoimmune disease or accidental damage. The polypeptides and nucleotides
CC may also be used in the diagnosis of the above conditions, and in drug
CC screening techniques. The present sequence represents a cDNA encoding a
CC novel human polypeptide of the invention
XX
SQ Sequence 5665 BP; 1077 A; 1634 C; 1687 G; 1267 T; 0 U; 0 Other;
Query Match 81.7%; Score 18.8; DB 4; Length 5665;
Best Local Similarity 90.9%; Pred. No. 1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 2 ATGTATTGATGGGATAGG 23
Db 1567 ATATATTGATGGGGAAGG 1588
|||||
RESULT 6
ADE81837
ID ADE81837 standard; cDNA; 493 BP.
XX
AC ADE81837;
XX
DT 29-JAN-2004 (first entry)
XX
DE Arabidopsis thaliana expressed polynucleotide seq id 608.
XX
KW genetically modified organism; transgenic organism; plant;
KW inhibitor testing; activator testing; modifier testing; fungicide;
KW insecticide; genetic function; genetic regulation; cellular metabolism;
KW gene; ss.
XX
OS Arabidopsis thaliana.
XX
PN US2003115639-A1.
XX
PD 19-JUN-2003.
XX
PF 26-JAN-2001; 2001US-00770961.
XX
PR 27-JAN-2000; 2000US-0178466P.
XX
PA (GORL/) GORLACH J.
PA (ANYV/) AN Y.
PA (HAML/) HAMILTON C M.
PA (PRIC/) PRICE J L.
PA (RAIN/) RAINES T M.
PA (YUYV/) YU Y.
PA (RAME/) RAMEAKA J G.
PA (PAGE/) PAGE A. V.
PA (MATH/) MATHAW A V.
PA (LEDF/) LEDFORD B L.
PA (WOES/) WOESSNER J P.
PA (HAAS/) HAAS W D.
PA (GARC/) GARCIA C A.
PA (KRIC/) KRICKER M.
PA (SLAT/) SLATER T.
PA (DAVI/) DAVIS K R.
PA (ALLE/) ALLEN K.
PA (HOFF/) HOFFMAN N.
PA (HURB/) HURBAN P.
XX

PR	19-JUL-1999	99US-01443333P
PR	19-JUL-1999	99US-01443334P
PR	19-JUL-1999	99US-01443335P
PR	20-JUL-1999	99US-01443352P
PR	20-JUL-1999	99US-0144632P
PR	20-JUL-1999	99US-0144884P
PR	21-JUL-1999	99US-0144814P
PR	21-JUL-1999	99US-0145086P
PR	21-JUL-1999	99US-0145088P
PR	22-JUL-1999	99US-0145085P
PR	22-JUL-1999	99US-0145087P
PR	22-JUL-1999	99US-0145089P
PR	22-JUL-1999	99US-01451192P
PR	23-JUL-1999	99US-01451145P
PR	23-JUL-1999	99US-0145218P
PR	23-JUL-1999	99US-0145224P
PR	26-JUL-1999	99US-0145276P
PR	26-JUL-1999	99US-01459133P
PR	27-JUL-1999	99US-0145918P
PR	27-JUL-1999	99US-0145919P
PR	28-JUL-1999	99US-0145951P
PR	28-AUG-1999	99US-0146386P
PR	02-AUG-1999	99US-0146388P
PR	02-AUG-1999	99US-0146389P
PR	03-AUG-1999	99US-0147038P
PR	04-AUG-1999	99US-0147204P
PR	04-AUG-1999	99US-0147302P
PR	05-AUG-1999	99US-0147192P
PR	05-AUG-1999	99US-0147260P
PR	06-AUG-1999	99US-0147303P
PR	06-AUG-1999	99US-0147416P
PR	09-AUG-1999	99US-0147493P
PR	09-AUG-1999	99US-01475935P
PR	10-AUG-1999	99US-0148171P
PR	10-AUG-1999	99US-0148319P
PR	12-AUG-1999	99US-0148341P
PR	13-AUG-1999	99US-0148565P
PR	13-AUG-1999	99US-0148684P
PR	16-AUG-1999	99US-0149368P
PR	17-AUG-1999	99US-0149175P
PR	18-AUG-1999	99US-0149426P
PR	20-AUG-1999	99US-0149722P
PR	20-AUG-1999	99US-0149723P
PR	20-AUG-1999	99US-0149929P
PR	23-AUG-1999	99US-0149502P
PR	23-AUG-1999	99US-0149930P
PR	25-AUG-1999	99US-0150566P
PR	26-AUG-1999	99US-0150884P
PR	27-AUG-1999	99US-0151065P
PR	27-AUG-1999	99US-0151066P
PR	30-AUG-1999	99US-0151303P
PR	31-AUG-1999	99US-0151438P
PR	01-SEP-1999	99US-0151930P
PR	07-SEP-1999	99US-0152363P
PR	10-SEP-1999	99US-0153070P
PR	13-SEP-1999	99US-0153758P
PR	15-SEP-1999	99US-0154018P
PR	20-SEP-1999	99US-0154039P
PR	22-SEP-1999	99US-0154177P
PR	23-SEP-1999	99US-0155139P
PR	24-SEP-1999	99US-0155486P
PR	28-SEP-1999	99US-0155659P
PR	29-SEP-1999	99US-0156458P
PR	04-OCT-1999	99US-0156596P
PR	05-OCT-1999	99US-0157117P
PR	06-OCT-1999	99US-0157753P
PR	07-OCT-1999	99US-0157865P
PR	08-OCT-1999	99US-0158029P
PR	12-OCT-1999	99US-0158232P
PR	13-OCT-1999	99US-0158369P
PR	13-OCT-1999	99US-0159293P
PR	13-OCT-1999	99US-0192924P

PR	13-OCT-1999;	99US-0159295P.
PR	14-OCT-1999;	99US-0159329P.
PR	14-OCT-1999;	99US-0159330P.
PR	14-OCT-1999;	99US-0159331P.
PR	14-OCT-1999;	99US-0159637P.
PR	14-OCT-1999;	99US-0159638P.
PR	18-OCT-1999;	99US-0159584P.
PR	21-OCT-1999;	99US-0160741P.
PR	21-OCT-1999;	99US-0160767P.
PR	21-OCT-1999;	99US-0160768P.
PR	21-OCT-1999;	99US-0160770P.
PR	21-OCT-1999;	99US-0160814P.
PR	22-OCT-1999;	99US-0160815P.
PR	22-OCT-1999;	99US-0160980P.
PR	22-OCT-1999;	99US-0160981P.
PR	22-OCT-1999;	99US-0160989P.
PR	25-OCT-1999;	99US-0161404P.
PR	25-OCT-1999;	99US-0161405P.
PR	25-OCT-1999;	99US-0161406P.
PR	26-OCT-1999;	99US-0161359P.
PR	26-OCT-1999;	99US-0161360P.
PR	26-OCT-1999;	99US-0161361P.
PR	28-OCT-1999;	99US-0161320P.
PR	28-OCT-1999;	99US-0161992P.
PR	28-OCT-1999;	99US-0161993P.
PR	29-OCT-1999;	99US-0162142P.

Query Match 80.0%; Score

Best Local Similarity 95.0%; Pred

Matches 19; Conservative 0; M

Qy	4	GTAATTGATGGGATAGAGG 23
Db	979	GCATTGATGGGATAGAGG 998

RESULT 8

ADRO6686

ID ADR06686 standard; cDNA; 3050 BP.

XX

AC ADR06686;

DT 04-NOV-2004 (first entry)

XX Full length human cDNA useful for

DE Gene; ss; human; oligo-capping me

XX osteoporosis; neurological diseas

KW Parkinson's disease; dementia; sh

KW sense or motor function; emotiona

KW osteopathic; neuroprotective; noo

KW transulliser.

XX

OS Homo sapiens.

XX

PX EP1447413-A2.

PN

XX

PD 18-AUG-2004.

XX

PF 12-FEB-2004; 2004EP-00003145.

XX

PR 14-FEB-2003; 2003JP-00102207.

PR 09-MAY-2003; 2003JP-00131452.

XX

XX (REAS-) RES ASSOC BIOTECHNOLOGY.

PA

XX Isoai T, Yamamoto J, Nishikawa

PI Wakamatsu A, Ishii S, Nagai K,

XX WPI; 2004-583265/57.

DR P-PSDB; ADR08642.

DR

XX New 1995 cDNA, useful for treatin

PT

PT Alzheimer's diseases, Parkinson's diseases, dementia and various cancers.
 XX Claim 1; SEQ ID NO 192; 2686pp; English.

XX This invention relates to novel, isolated full length human cDNA
 CC molecules and the encoded proteins thereof. Specifically, it refers to
 CC cDNA clones obtained by an oligo-capping method, where none of these
 CC clones are identical to any known human mRNAs. The present invention
 CC describes an immunoassay to identify agonists and antagonists, as well as
 CC antibodies, antisense molecules and siRNAs that can all be used to bind
 CC to and modulate expression of the cDNA molecules. As such, these
 CC molecules are useful for diagnostic markers or therapeutic targets for
 CC the various diseases or morbid states. In particular, they are useful in
 CC gene therapy for treating osteoporosis, neurological disease, Alzheimer's
 CC disease, Parkinson's disease, dementia, short memory and various cancers,
 CC as well as for maintaining equilibrium of sense or motor function, and
 CC for treating emotional reaction, fear response and panic. Accordingly,
 CC they exhibit osteoprotective, neuroprotective, nootropic, antiparkinsonian,
 CC cyostatic and tranquiliser activities. This polynucleotide is a full
 CC length human cDNA sequence of the invention. NOTE: This sequence is not
 CC given in the sequence listing of the specification but can be obtained on
 CC CD-ROM from the European Patent Office, Vienna Sub-office.

XX SQ Sequence 3050 BP; 978 A; 562 C; 645 G; 865 T; 0 U; 0 Other;

Query Match 80.0%; Score 18.4; DB 13; Length 3050;
 Best Local Similarity 95.0%; Pred. No. 1.5e+02;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 ATGTATTGATGGGATAGA 21
 |||||
 Db 1616 ATGTATTGATGGGATACA 1635

RESULT 9

ACC00653
 ID ACC00653 standard; cDNA; 1630 BP.

XX AC ACC00653;

XX DT 16-MAY-2003 (first entry)

XX DE Argemone mexicana oil trait related cDNA sequence SEQ ID NO:55.

XX KW Plant; oil trait; oil phenotype; altered lipid profile; MAP kinase;
 KW receptor-like protein kinase; mitogen activated protein kinase; oil;
 KW Lipid biosynthesis; Hap2-like; Haps-like; Hap3/Lec1-like; prickly poppy.
 KW CKC-like transcription factor; calcosin; ATP citrate lyase; SNF1;
 KW transgenic plant; gene; ss.

XX OS Argemone mexicana.

XX PN WO2003002751-A2.

XX PD 09-JAN-2003.

XX PF 27-JUN-2002; 2002WO-US020152.

XX PR 29-JUN-2001; 2001US-0301913P.

XX PA (DUPO) DU PONT DE NEMOURS & CO E I.
 PA (PION-) PIONEER HI-BRED INT INC.

XX PI Allen SM, Allen WB, Cahoon RE, Epelbaum S, Famodu OO, Harvell LT;
 PI Jones TJ, Kinney AJ, Klein TM, Li C, Oliveira IC, Sakai H, Shen B;
 PI Tarczynski MC;

XX DR WPI; 2003-201509/19.
 DR P-PSDB; ABR40616.

XX Novel nucleotide fragment encoding polypeptides having receptor-like
 PT protein kinase activity, calcosin-like activity, useful for altering oil
 PT phenotypes in plants such as sunflower, coconut, soybean, wheat and rice.

XX Claim 18; Page 129; 542pp; English.

XX The present invention describes an isolated nucleotide fragment (I)
 CC comprising a nucleic acid sequence (NS) chosen from a NS encoding a
 CC polypeptide (PP) having receptor-like protein kinase activity, mitogen
 CC activated protein (MAP)-kinase activity, Lipid-like transcription factor
 CC activity, calcosin-like activity, ATP citrate lyase activity, SNF1-like
 CC activity and CKC-like transcription factor activity. Also described: (1)
 CC complement (II) of (I); (2) a chimeric construct (III) comprising (I) or
 CC (II), operably linked to a regulatory sequence; (3) a plant (IV)
 CC comprising (III) in its genome; (4) seeds (V) obtained from (IV); and (5)
 CC oil obtained from (V). (I) or its part can be used in antisense
 CC inhibiting or co-suppression in a transformed plant. (III) is useful for
 CC altering the oil phenotype in a plant such as corn, soybean, wheat, rice,
 CC canola, Brassica, sorghum, sunflower or coconut. (III) is also useful for
 CC creating transgenic plants having altered lipid profiles. (I) can also be
 CC used as a hybridisation probe. ACC00626 to ACC00868 and ABR40591 to
 CC ABR40879 represent sequences used in the exemplification of the present
 CC invention

XX SQ Sequence 1630 BP; 497 A; 318 C; 375 G; 440 T; 0 U; 0 Other;

Query Match 79.1%; Score 18.2; DB 8; Length 1630;
 Best Local Similarity 87.0%; Pred. No. 1.7e+02;
 Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CATGTATTGATGGGATAGAGG 23
 |||||
 Db 1328 CTTGTGTTTGTGGGAAAGAGG 1350

RESULT 10

ADC23534

ID ADC23534 standard; cDNA; 1630 BP.

XX AC ADC23534;

XX DT 18-DEC-2003 (first entry)

XX DE cDNA encodes protein involved in altering plant oil phenotype (SeqID 39).

XX ss; gene; oil phenotype; plant; breeding; Hap transcription factor;
 KW lipid biosynthesis; Hap2-like; Haps-like; Hap3/Lec1-like; prickly poppy.

XX OS Argemone mexicana.

XX PN WO2003001902-A2.

XX PD 09-JAN-2003.

XX PF 27-JUN-2002; 2002WO-US022086.

XX PR 29-JUN-2001; 2001US-0301913P.

XX PA (DUPO) DU PONT DE NEMOURS & CO E I.
 PA (PION-) PIONEER HI-BRED INT INC.

XX PI Allen WB, Cahoon RE, Famodu OO, Harvell LT, Helentjaris TG, Li C;
 PI Lowe KS, Oliveira IC, Shen B, Tarczynski MC;
 PI WPI; 2003-210187/20.

XX DR P-PSDB; ADC23535.

XX Novel nucleotide fragment encoding polypeptides having Hap2, Hap5 or
 PT Hap3/Lec1-like activity useful for altering oil phenotypes in plants such
 PT as sunflower, coconut, soybean, wheat and rice.

XX Claim 19; SEQ ID NO 39; 202pp; English.

XX The present invention relates to a novel method for altering oil
 CC phenotype in plants through the controlled expression of selective genes,
 CC useful in plant breeding. Specifically, it refers to the isolated nucleic

AC ACN44424;
 XX
 DT 18-NOV-2004 (first entry)
 XX
 DE Mouse genomic sequence MCG19046.
 XX
 KW Cytostatic; carcinoma; lymphoma; cancer; murine; gene; ss.
 XX
 OS Mus musculus.
 XX
 XX WC2003073826-A2.
 XX
 PD 12-SEP-2003.
 XX
 PF 28-FEB-2003; 2003WO-US006235.
 XX
 XX 01-MAR-2002; 2002US-00087192.
 XX
 PA (SAGR-) SAGRES DISCOVERY.
 XX
 PI Morris DW;
 XX
 XX WPI; 2003-328604/31.
 XX
 PT Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
 PT comprises a nucleotide sequence.
 XX
 PS Claim 1; SEQ ID NO 865; Opp; English.
 XX
 CC The present invention relates to novel DNA and protein sequences which
 CC are associated with carcinomas. The sequences are useful for: (i) for
 CC screening drug candidates; (ii) for screening of bioactive agent capable
 CC of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
 CC a bioactive agent capable of modulating the activity of CAP; (iv) for
 CC evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
 CC carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
 CC carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
 CC (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
 CC determining Carcinoma Associated (CA) gene copy number. In addition, the
 CC CA genes are useful as DNA vaccines and the CAP are useful as markers of
 CC carcinoma including lymphoma. The present sequence is one such CA coding
 CC sequence. Note: This patent is an equivalent to basic patent
 CC US2002182586A1, for which no sequence data was published
 XX
 SQ Sequence 29111 BP; 6425 A; 7887 C; 7445 G; 6494 T; 0 U; 860 Other;
 Query Match 79.1%; Score 18.2; DB 11; Length 29111;
 Best Local Similarity 79.0%; Pred. No. 2.5e+02;
 Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 QY 1 CATGTATTTGATGGGATAGG 23
 DB 22398 CATGTATGGATGGGAGAGAGG 22376
 RESULT 14
 ID ADS46987/c
 XX ADS46987 standard; cDNA; 2286 BP.
 XX
 AC ADS46987;
 XX
 XX 02-DEC-2004 (first entry)
 XX
 XX Bacterial polynucleotide #1730.
 XX
 KW Recombinant DNA construct; transformed plant; improved plant property;
 KW cold tolerance; heat tolerance; drought tolerance; herbicide; osmosis;
 KW pathogen tolerance; pest tolerance; plant disease resistance;
 KW cell cycle pathway modification; plant growth regulator;
 KW homologous recombination; seed oil yield; protein yield; carbohydrate;
 KW nitrogen; phosphorus; photosynthesis; lignin; galactomannan;
 KW bacterial polynucleotide; gene; ss.
 XX

OS Bacteria.
 XX
 PN US2003233675-A1.
 XX
 PD 18-DEC-2003.
 XX
 XX 20-FEB-2003; 2003US-00369493.
 PF
 PR 21-FEB-2002; 2002US-0360039P.
 XX
 XX (CAOY/) CAO Y.
 PA (HINK/) HINKLE G J.
 PA (SLAY/) SLATER S C.
 PA (CHEN/) CHEN X.
 PA (GOLD/) GOLDMAN B S.
 XX
 PI Cao Y, Hinkle GJ, Slater SC, Chen X, Goldman BS;
 XX
 XX WPI; 2004-061375/06.
 XX
 PT New recombinant DNA construct comprising a promoter positioned to provide
 PT for expression of a polynucleotide encoding a polypeptide from a
 PT microbial source, useful for producing plants with improved properties.
 XX
 PS Claim 1; SEQ ID NO 25417; 122pp; English.
 XX
 CC The invention relates to a recombinant DNA construct comprising a
 CC promoter functional in a plant cell, where the promoter is positioned to
 CC provide for expression of a polynucleotide encoding a polypeptide from a
 CC microbial source. The invention also relates to a transformed plant
 CC comprising the recombinant DNA construct and a method of producing a
 CC transformed plant having an improved property. The plant is a crop plant
 CC such as maize or soybean. The method of producing a transformed plant
 CC having an improved property comprises transforming a plant with the
 CC recombinant DNA construct and growing the transformed plant, where the
 CC polynucleotide or polypeptide is useful for improving plant properties.
 CC The recombinant DNA construct is useful for producing plants with
 CC tolerance plant properties, e.g. improved cold, heat or drought tolerance,
 CC tolerance to herbicides, extreme osmotic conditions, pathogens or pests,
 CC increased resistance to plant disease, better growth rate by modification
 CC of the cell cycle pathway with plant growth regulators, increased rate of
 CC homologous recombination, modified seed oil or protein yield and/or
 CC content, improved yield by modification of carbohydrate, nitrogen or
 CC phosphorus use and/or uptake, by modification of photosynthesis or by
 CC providing improved plant growth and development under at least one stress
 CC condition, improved lignin production or improved galactomannan
 CC production. This sequence represents a bacterial polynucleotide used in
 CC the scope of the invention. Note: The sequence data for this patent did
 CC not form part of the printed specification but was obtained in electronic
 CC format from USPTO at seqdata.uspto.gov/sequence.html.
 XX
 SQ Sequence 2286 BP; 697 A; 474 C; 512 G; 603 T; 0 U; 0 Other;
 Query Match 78.3%; Score 18; DB 13; Length 2286;
 Best Local Similarity 100.0%; Pred. No. 2.2e+02;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 ATGTATTTCATGGGATA 19
 DB 656 ATGTATTTCATGGGATA 639
 RESULT 15
 ID ABV12937/c
 XX ABV12937 standard; cDNA; 238 BP.
 XX
 AC ABV12937;
 XX
 XX 13-SEP-2002 (first entry)
 XX
 DE Human prostate expression marker cDNA 12928.
 XX
 KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynanamic marker;

KW pharmacogenomic marker; gene; ss.
XX
OS Homo sapiens.
XX
FN WO200160860-A2.
XX
XX 23-AUG-2001.
XX
PD
XX
PF 20-FEB-2001; 2001WO-US005171.
XX
XX 17-FEB-2000; 2000US-0183319P.
PR 16-MAR-2000; 2000US-0189862P.
PR 25-MAY-2000; 2000US-0207454P.
PR 09-JUN-2000; 2000US-0211314P.
PR 18-JUL-2000; 2000US-0219007P.
PR 13-DEC-2000; 2000US-0255281P.
XX
PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
XX
XX Schlegel R, Endege WO, Monahan JE;
PI
XX WPI; 2001-662795/76.
XX
XX Novel isolated nucleic acid molecule associated with cancerous state of
PT prostate cells and correlating with presence of prostate cancer, useful
PT for detecting presence of prostate cancer, stage of prostate cancer.
XX
PS Claim 1; Page 2134; 11750pp; English.
XX
XX The invention relates to an isolated nucleic acid molecule (I) comprising
CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the
CC specification or its complement. (I) is useful for: (a) assessing whether
CC a patient is afflicted with prostate cancer; (b) monitoring the
CC progression of prostate cancer in a patient; (c) assessing the efficacy
CC of a test compound to inhibit prostate cancer in a patient; (d) assessing
CC the efficacy of a therapy for inhibiting prostate cancer in a patient;
CC (e) selecting a composition for inhibiting prostate cancer in a patient;
CC (f) assessing the prostate cell carcinogenic potential of a compound; (g)
CC determining whether prostate cancer has metastasized in a patient; (h)
CC assessing the aggressiveness or indolence of prostate cancer in a patient
CC ; (I) is also useful as a pharmacodynamic or pharmacogenomic marker
XX
SQ Sequence 238 BP; 79 A; 60 C; 36 G; 63 T; 0 U; 0 Other;

Query Match 77.4%; Score 17.8; DB 5; Length 238;
Best Local Similarity 90.5%; Pred. No. 2e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TGTATTGTGGGGATAGGG 23
Db 54 TGTATTGTGGGGATAGGG 34

Search completed: August 13, 2005, 04:14:28
Job time : 190.245 secs

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		Match	Length			
1	18.8	81.7	93364	4	US-09-949-016-14890	Sequence 14890, A
2	17.8	77.4	89689	4	US-09-949-016-13089	Sequence 13089, A
3	17.8	77.4	157644	4	US-09-949-016-16179	Sequence 16179, A
4	17.2	77.4	157644	4	US-09-949-016-16180	Sequence 16180, A
5	17.2	74.8	1026	4	US-09-710-279-1277	Sequence 1277, A
6	17.2	74.8	2250	4	US-09-710-279-137	Sequence 137, App
7	17.2	74.8	2448	3	US-09-134-001C-777	Sequence 777, App
8	17.2	74.8	3427	4	US-09-710-279-4289	Sequence 4289, App
9	17.2	74.8	4198	4	US-09-710-279-3604	Sequence 3604, App
10	16.8	73.0	71251	4	US-09-949-016-15332	Sequence 15332, A
11	16.6	72.2	209	4	US-09-313-294A-765	Sequence 765, App
12	16.6	72.2	211	4	US-09-513-999C-32576	Sequence 32576, A
13	16.6	72.2	601	4	US-09-949-016-64747	Sequence 64747, A
14	16.6	72.2	827	3	US-09-328-111-93	Sequence 93, Appl
15	16.6	72.2	3880	3	US-09-221-017B-1071	Sequence 1071, App
16	16.6	72.2	25274	4	US-09-949-016-16682	Sequence 16682, A
17	16.6	72.2	57751	4	US-09-949-016-13631	Sequence 13631, A
18	16.4	71.3	1917	4	US-09-134-000C-1459	Sequence 1459, App
19	16.4	71.3	103712	4	US-09-949-016-13058	Sequence 13058, A
20	16.4	71.3	236341	4	US-09-949-016-13978	Sequence 13978, A
21	16.2	70.4	277	4	US-09-313-294A-130	Sequence 130, App
22	16.2	70.4	307	4	US-09-313-294A-6619	Sequence 6619, App
23	16.2	70.4	441	4	US-09-107-532A-2725	Sequence 2725, App
24	16.2	70.4	601	4	US-09-949-016-205970	Sequence 205970, A
25	16.2	70.4	912	4	US-09-107-532A-833	Sequence 833, App
26	16.2	70.4	1941	4	US-09-543-681A-2668	Sequence 2668, App
27	16.2	70.4	2250	4	US-09-902-540-5197	Sequence 5197, App

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; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13089
; LENGTH: 89689
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(89689)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13089

Query Match      77.4%; Score 17.8; DB 4; Length 89689;
Best Local Similarity 90.5%; Pred. No. 1.7e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CATGTATTTCATGGGATAG 21
Db 14881 CATGTATTTCATGGGACAGA 14901

RESULT 3
US-09-949-016-16179
; Sequence 16179, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16179
; LENGTH: 157644
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(157644)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-16179

Query Match      77.4%; Score 17.8; DB 4; Length 157644;
Best Local Similarity 90.5%; Pred. No. 1.7e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTATTTCATGGGATAG 22
Db 150480 ATGTATTTCATGGGCTAG 150500

RESULT 4
US-09-949-016-16180
; Sequence 16180, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
```

```
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16180
; LENGTH: 157644
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(157644)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-16180

Query Match      77.4%; Score 17.8; DB 4; Length 157644;
Best Local Similarity 90.5%; Pred. No. 1.7e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTATTTCATGGGATAG 22
Db 150480 ATGTATTTCATGGGCTAG 150500

RESULT 5
US-09-710-279-1277
; Sequence 1277, Application US/09710279
; Patent No. 6703492
; GENERAL INFORMATION:
; APPLICANT: KIMMERLY, WILLIAM JOHN
; TITLE OF INVENTION: STAPHYLOCOCCUS EPIDERMIDIS NUCLEIC ACIDS AND PROTEINS
; FILE REFERENCE: PU3480US
; CURRENT APPLICATION NUMBER: US/09/710,279
; CURRENT FILING DATE: 2000-11-09
; PRIOR APPLICATION NUMBER: 60/164,258
; PRIOR FILING DATE: 1999-11-09
; NUMBER OF SEQ ID NOS: 4472
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1277
; LENGTH: 1026
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: synthetic
; OTHER INFORMATION: nucleic acid sequence
US-09-710-279-1277

Query Match      74.8%; Score 17.2; DB 4; Length 1026;
Best Local Similarity 86.4%; Pred. No. 1.6e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 ATGTATTTCATGGGATAG 23
Db 470 ATATATTTAATGGGATATAG 491

RESULT 6
US-09-710-279-137
; Sequence 137, Application US/09710279
; Patent No. 6703492
; GENERAL INFORMATION:
; APPLICANT: KIMMERLY, WILLIAM JOHN
; TITLE OF INVENTION: STAPHYLOCOCCUS EPIDERMIDIS NUCLEIC ACIDS AND PROTEINS
; FILE REFERENCE: PU3480US
; CURRENT APPLICATION NUMBER: US/09/710,279
; CURRENT FILING DATE: 2000-11-09
; PRIOR APPLICATION NUMBER: 60/164,258
; PRIOR FILING DATE: 1999-11-09
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US-09-710-279-4289

Db 51291 TGTATTGTGGGAGAG 51310

RESULT 11

US-09-313-294A-765
; Sequence 765, Application US/09313294A
; Patent No. 6476212
; GENERAL INFORMATION:
; APPLICANT: Lalgudi, Raghunath V.
; APPLICANT: Ito, Laura Y.
; APPLICANT: Sherman, Bradley K.
; TITLE OF INVENTION: POLYNUCLEOTIDES AND POLYPEPTIDES DERIVED FROM CORN EAR
; FILE REFERENCE: PL-0017 US
; CURRENT APPLICATION NUMBER: US/09/313,294A
; CURRENT FILING DATE: 1999-05-14
; NUMBER OF SEQ ID NOS: 760
; SOFTWARE: PERL Program
; SEQ ID NO 765
; LENGTH: 209
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. 6476212 700549795H1
; NAME/KEY: unsure
; LOCATION: 29, 37, 39, 52, 57, 71, 76, 85, 118, 153-154, 201
; OTHER INFORMATION: a, t, c, g, or other
US-09-313-294A-765

Query Match 72.2%; Score 16.6; DB 4; Length 209;

Best Local Similarity 82.6%; Pred. No. 2.4e+02; Mismatches 0; Indels 0; Gaps 0;

QY 1 CATGTATTGTGGGATAGG 23

Db 4 CATGTATCTGGTGGATGTGG 26

RESULT 12

US-09-513-999C-22576
; Sequence 22576, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59 US2 REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 22576
; LENGTH: 211
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 40
; OTHER INFORMATION: r=a or g
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 184
; OTHER INFORMATION: k=g or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 202
; OTHER INFORMATION: k=g or t
US-09-513-999C-22576

Query Match 72.2%; Score 16.6; DB 4; Length 211;
Best Local Similarity 82.6%; Pred. No. 2.4e+02; Mismatches 0; Indels 0; Gaps 0;

QY 1 CATGTATTGTGGGATAGG 23

Db 72 CATGAATTTGGGATATGG 94

RESULT 13

US-09-949-016-64747
; Sequence 64747, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 64747
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-64747

Query Match 72.2%; Score 16.6; DB 4; Length 601;

Best Local Similarity 82.6%; Pred. No. 2.8e+02; Mismatches 0; Indels 0; Gaps 0;

QY 1 CATGTATTGTGGGATAGG 23

Db 189 CATGTGTGTGGTGAGAGG 211

RESULT 14

US-09-328-111-93
; Sequence 93, Application US/09328111
; Patent No. 6262333
; GENERAL INFORMATION:
; APPLICANT: Endege, Wilson O.
; APPLICANT: Steinmann, Kathleen E.
; APPLICANT: Astle, Jon H.
; APPLICANT: Burgess, Christopher C.
; APPLICANT: Bushnell, Steven E.
; APPLICANT: Carroll III, Eddie
; APPLICANT: Catino, Theodore J.
; APPLICANT: Derti, Adnan
; APPLICANT: Ford, Donna M.
; APPLICANT: Lewis, Marcia E.
; APPLICANT: Monahan, John E.
; APPLICANT: Schlegel, Robert
; TITLE OF INVENTION: NOVEL HUMAN GENES AND GENE EXPRESSION
; FILE REFERENCE: CCD-257 (US)
; CURRENT APPLICATION NUMBER: US/09/328,111
; CURRENT FILING DATE: 1999-06-08
; EARLIER APPLICATION NUMBER: US 60/088,801
; EARLIER FILING DATE: 1998-06-10
; NUMBER OF SEQ ID NOS: 850
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 93
; LENGTH: 627
; TYPE: DNA
; ORGANISM: Homo sapiens

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: August 13, 2005, 03:26:33 ; Search time 1478.12 Seconds
(without alignments)
592.293 Million cell updates/sec

Title: US-10-673-854-1

Perfect score: 23

Sequence: 1 catgtatttgatgggatagag 23

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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6: gb_est5.*

7: gb_est6.*

8: gb_gss1.*

9: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 2	18.8	81.7	401	1	AA445156 vf58f07.y
C 3	18.8	81.7	570	8	AZ662915 1M0542H17
C 4	18.8	81.7	616	5	BQ829481 L6in2040
C 5	18.8	81.7	659	9	CE717779 tigr-gss-
C 6	18.8	81.7	685	9	CC514760 CH240 359
C 7	18.8	81.7	946	2	BF580964 602100666
C 8	18.8	81.7	952	5	BQ929148 AGENCOURT
C 9	18.8	81.7	979	5	BQ941451 AGENCOURT
C 10	18.8	81.7	987	8	CC204478 CH261-131
C 11	18.8	81.7	989	4	BG296600 602393522
C 12	18.4	80.0	323	8	BH854982 SALK 0871
C 13	18.4	80.0	365	7	T75599 10377 Lambd
C 14	18.4	80.0	372	5	BP658944 BP658944
C 15	18.4	80.0	400	1	AV814465 AV814465
C 16	18.4	80.0	404	8	B24975 F23124TF IG
C 17	18.4	80.0	406	1	AV786094 AV786094
C 18	18.4	80.0	406	1	AV806091 AV806091
C 19	18.4	80.0	409	5	BP662526 BP662526
C 20	18.4	80.0	415	1	AV786678 AV786678
C 21	18.4	80.0	452	6	CD529428 02N16 Ara
C 22	18.4	80.0	452	9	AL087865 Arabidops
C 23	18.4	80.0	458	1	AI928235 701493742
C 24	18.4	80.0	483	1	AA586206 28855 Lam

25	18.4	80.0	525	9	TA288A02P	AL487137 T. brucei
C 26	18.4	80.0	556	9	TA59G11P	AL455675 T. brucei
27	18.4	80.0	585	9	TA189G12Q	AL476670 T. brucei
28	18.4	80.0	908	7	CK412987	CK412987 AUF IpGill
29	18.4	80.0	1214	3	CNS0A6GM	BX824231 Arabidops
30	18.4	80.0	1227	3	CNS0A4TV	BX85041 Arabidops
31	18.4	80.0	1271	3	CNS0A77H	BX822669 Arabidops
32	18.4	80.0	1293	3	CNS0A6JM	BX823458 Arabidops
33	18.4	80.0	1302	3	CNS0A6IY	BX823867 Arabidops
34	18.2	79.1	195	2	BB584673	BB584673 Arabidops
C 35	18.2	79.1	306	1	AV204432	AV204432 Arabidops
C 36	18.2	79.1	474	2	BP869304	BP869304 IL3-ET011
C 37	18.2	79.1	538	8	AQ143748	AQ143748 HS_3075_B
38	18.2	79.1	589	7	CR791845	CR791845 DKF2P468N
39	18.2	79.1	611	7	CK943116	CK943116 4067003 B
40	18.2	79.1	654	6	CF073177	CF073177 FE1 28 G0
41	18.2	79.1	674	8	AQ040543	AQ040543 CIT-HSP-2
C 42	18.2	79.1	687	9	CE737028	CE737028 tigr-gss-
C 43	18.2	79.1	713	8	AZ246641	AZ246641 RPCI-23-B
C 44	18.2	79.1	736	7	CK943520	CK943520 4067387 B
C 45	18.2	79.1	747	9	CL839877	CL839877 OR_Cba006

ALIGNMENTS

RESULT 1	AI552179/c	AI552179	394 bp	mrna	linear	EST 23-MAR-1999
LOCUS	VF58F07.Y1	Soares mouse NBMH Mus musculus cDNA clone IMAGE:848005				
DEFINITION	5' similar to gb:U07592 PEROXISOME PROLIFERATOR ACTIVATED RECEPTOR BETA (HUMAN); mRNA sequence.					
ACCESSION	AI552179	AI552179.1	GI:4484553			
VERSION	EST.					
KEYWORDS	Mus musculus (house mouse)					
SOURCE	Mus musculus					
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.					
REFERENCE	1 (bases 1 to 394)					
AUTHORS	Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T., Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person, B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk, R., Ritter, E., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and Wilson, R.					
TITLE	The WashU-NCI Mouse EST Project 1999					
JOURNAL	Unpublished (1999)					
COMMENT	Contact: Marra M/WashU-NCI Mouse EST Project 1999 Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800 Fax: 314 286 1810 Email: mouseest@watson.wustl.edu This clone is available royalty-free through LNL ; contact the IMAGE Consortium (info@image.llnl.gov) for further information. MGI:500157 This read is a RESEQUENCE of a previously sequenced mouse clone This read has been verified (found to hit its original self in the correct orientation) Seq primer: -40RP from Gibco High quality sequence stop: 382.					
FEATURES	Location/Qualifiers					
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	/mol_type="mRNA"					
	/strain="C57BL/6J"					
	/db_xref="taxon:10090"					
	/clone="IMAGE:848005"					
	/sex="male"					
	/tissue_type="heart"					
	/dev_stage="4 weeks"					
	/lab_host="DH10B"					
	/clone_lib="Soares mouse NBMH"					

/note="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' TGTTACCAATCGAAGTGGAGCGCGCGAAGTTTTTTTTTTTTTTTTTTT 3']; double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. RNA provided by Dr. Minoru Ko, Wayne State Univ. Library constructed and normalized by Bento Soares and M.Fatima Bonaldo."

ORIGIN

Query Match 81.7%; Score 18.8; DB 1; Length 394;
Best Local Similarity 90.9%; Pred. No. 6.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CATGTATTGATGGGGATAG 22
||||| ||||| ||||| |||||
Db 164 CATGCTTTTGAAGGGATAG 143

RESULT 2

AA445156/c
LOCUS
DEFINITION
v59f07.r1 Soares mouse NbMH Mus musculus cDNA clone IMAGE:848005
5' similar to gb:L07592 PEROXISOME PROLIFERATOR ACTIVATED RECEPTOR
BETA (HUMAN); mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
AA445156
AA445156.1 GI:2157839
Mus musculus (house mouse)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 401)

REFERENCE

AUTHORS
Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
Schellenberg, K., Steptoe, M., Tan, P., Underwood, K., Moore, B.,
Theising, B., Wyllie, T., Lennon, G., Soares, B., Wilson, R. and
Waterston, R.

TITLE

The WashU-HHMI Mouse EST Project

JOURNAL

Unpublished (1996)

COMMENT

Contact: Marra M/Mouse EST Project
WashU-HHMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

Email: mouseest@watson.wustl.edu

This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.

MG1:500157

Seq primer: -28ml3 rev2 ET from Amersham

High quality sequence stop: 385.

FEATURES

source

1..401
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="IMAGE:848005"
/sex="male"
/tissue_type="heart"
/dev_stage="4 weeks"
/lab_host="DH10B"
/clone_lib="Soares mouse NbMH"

/note="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' TGTTACCAATCGAAGTGGAGCGCGCGAAGTTTTTTTTTTTTTTTTTTT 3']; double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. RNA

provided by Dr. Minoru Ko, Wayne State Univ. Library
constructed and normalized by Bento Soares and M.Fatima
Bonaldo."

ORIGIN

Query Match 81.7%; Score 18.8; DB 1; Length 401;
Best Local Similarity 90.9%; Pred. No. 6.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CATGTATTGATGGGGATAG 22
||||| ||||| ||||| |||||
Db 158 CATGCTTTTGAAGGGATAG 137

RESULT 3

AZ662915
LOCUS
DEFINITION
1M0542H17F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0542H17 F, genomic survey sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
AZ662915
AZ662915.1 GI:11800061
Mus musculus (house mouse)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 570)

REFERENCE

AUTHORS
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T.,
Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von
Niederhausen, A. and Wright, D., Weiss, R.

Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

JOURNAL

Unpublished (2000)

COMMENT

Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0542 row: H column: 17

Seq primer: CTTGTAAACGACGCCAGT

Class: plasmid ends

High quality sequence stop: 570.

FEATURES

Location/Qualifiers

1..570

/organism="Mus musculus"

/mol_type="genomic DNA"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC1M0542H17"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"

/clone_lib="Mouse 10kb plasmid UUGC1M library"

/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 [gi|4732114|gb|AF129072.1], a copy-number
inducible derivative of plasmid RI. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to

adapted vector DNA, and transformed into chemically-competent *E. coli* XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

ORIGIN
Query Match 81.7%; Score 18.8; DB 8; Length 570;
Best Local Similarity 90.9%; Pred. No. 7.1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTATTGATGGGATAGAGG 23
|||||
Db 502 ATGTATTGTTGAGATAGAGG 523
|||||

RESULT 4
BQ829481/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
COMMENT

BQ829481 616 bp mRNA linear EST 15-SEP-2002
LL6in20408 AFT024-subtracted library Mus musculus cDNA 5' similar
to peroxisome proliferator activated receptor delta, mRNA sequence.
BQ829481
BQ829481.1 GI:22861536
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 616)
Lemischka, I.R. and Moore, K.A.
Hackney, J.A., Charbord, P., Brunk, B.P., Stoeckert, C.J.,
A molecular profile of a hematopoietic stem cell niche
Proc. Natl. Acad. Sci. U.S.A. 99 (20), 13061-13066 (2002)
12247628
12226475
Contact: Moore, Kateri A.
Department of Molecular Biology
Princeton University
217 Lewis Thomas Laboratory, Washington Road, Princeton, NJ 08544,
USA
Tel: 609 258 0605
Fax: 609 258 2759
Email: kmoore@molbio.princeton.edu
These ESTs are derived from a subtracted cDNA library enriched for
gene products expressed by a hematopoietic stem cell-supporting
stromal cell line, AFT024.
Seq primer: M13Reverse or T7.

FEATURES
source
1..616
Location/Qualifiers

/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/tissue_type="Fetal Liver"
/cell_type="Stromal cell"
/cell_line="AFT024"
/dev_stage="Embryonic day 14-14.5"
/lab_host="DH10B"
/clone_lib="AFT024-subtracted library"
/note="Organ: Fetal Liver; Vector: Sport 1; Site 1: Sal I;
Site 2: Not I; Two directionally cloned cDNA libraries
were made from a hematopoietic stem cell-supporting
stromal cell line (AFT024) and from a non-supporting
stromal cell line (2018). Subtractive hybridization was
performed by hybridization of the target, AFT024, single
stranded cDNA library in pSport1 to biotinylated RNA
transcribed from the driver, 2018 cDNA library in pSport2
with inserts cloned in the complementary orientation. The
AFT024-subtracted library contains 4.2x105 clones and is
depleted of common housekeeping gene products eg.
beta-actin and enriched for transcripts specific to
AFT024. For detailed protocols and additional information
please see our website at
<http://stromalcell.princeton.edu>."

ORIGIN

Query Match 81.7%; Score 18.8; DB 5; Length 616;
Best Local Similarity 90.9%; Pred. No. 7.2e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CATGTATTGATGGGATAGAG 22
|||||
Db 284 CATGCTTTTGAAGGGATAGAG 263
|||||

RESULT 5
CE717779
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
COMMENT

CE717779 659 bp DNA linear GSS 29-SEP-2003
tigr-ges-dog-17000314586051 Dog Library Canis familiaris genomic,
genomic survey sequence.
CE717779
CE717779.1 GI:37037217
GSS.
Canis familiaris (dog)
Canis familiaris
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
1 (bases 1 to 659)
Kirkness, E.F., Bafna, V., Halpern, A.L., Levy, S., Remington, K.,
Rusch, D.B., Delcher, A.L., Pop, M., Wang, W., Fraser, C.M. and
Venter, J.C.
The dog genome: survey sequencing and comparative analysis
Science 301 (5641), 1898-1903 (2003)
22875432
14512627
Contact: Kirkness EF
The Institute for Genomic Research
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
Rockville, MD 20850, USA
Tel: 301-838-0200
Fax: 301-838-0208
Email: ekirknes@tigr.org
Class: shotgun.

FEATURES
source
1..659
Location/Qualifiers

/organism="Canis familiaris"
/mol_type="genomic DNA"
/strain="Standard Poodle"
/db_xref="taxon:9615"
/clone_lib="Dog Library"
/note="Site 1: BstXI; Libraries were prepared from
peripheral blood"

ORIGIN

Query Match 81.7%; Score 18.8; DB 9; Length 659;
Best Local Similarity 90.9%; Pred. No. 7.2e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTATTGATGGGATAGAGG 23
|||||
Db 571 ATGTTTTCATGGGATACAGG 592
|||||

RESULT 6
CC514760/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS

CC514760 685 bp DNA linear GSS 17-JUN-2003
CH240_359D20.T7 CHORI-240 Bos taurus genomic clone CH240_359D20,
genomic survey sequence.
CC514760
CC514760.1 GI:31833048
GSS.
Bos taurus (cow)
Bos taurus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovinae; Bos.
1 (bases 1 to 685)
Holt, R., Stott, J., Yang, G., Barber, S., Smalil, D., Prabhu, A.-L.,


```

QY 1 CATGTATTGATGGGGATAGAG 22
Db 576 CATGCTCTTGAAGGGGATAGAG 555

RESULT 9
BQ941451/c
LOCUS
DEFINITION AGENCOURT 8819379 NIH_MGC_18 Homo sapiens cDNA clone IMAGE:6422948
5', mRNA sequence.
ACCESSION BQ941451
VERSION BQ941451.1 GI:22356929
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE BQ941451
AUTHORS AGENCOURT
TITLE NIH-MGC http://mgc.nci.nih.gov/.
JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)
COMMENT Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgabbs-remail.nih.gov
Tissue Procurement: DCTD/DTP/Gazdar
cDNA Library Preparation: Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: L1CM2600 row: 1 column: 21
High quality sequence start: 15
High quality sequence stop: 596.

FEATURES
source
1..979
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6422948"
/tissue_type="large cell carcinoma"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH_MGC_18"
/notes="Organ: lung; Vector: pOTB7; Site 1: XhoI; Site 2:
EcoRI; cDNA made by oligo-dt priming. Directionally cloned
into EcoRI/XhoI sites using the following 5' adaptor:
GGCACAG(G). Library constructed by Ling Hong in the
laboratory of Gerald M. Rubin (University of California,
Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and
Superscript II RT (Life Technologies). Note: this is a
NIH_MGC Library."

ORIGIN
Query Match 81.7%; Score 18.8; DB 5; Length 979;
Best Local Similarity 90.9%; Pred. No. 7.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATGTATTGATGGGGATAGAG 23
Db 864 ATGTATTGATGGGGTTATAGG 843

RESULT 10
CC204478/c
LOCUS
DEFINITION CC204478 Sp6.1 CH261 Gallus gallus genomic clone CH261-131C12,
genomic survey sequence.
ACCESSION CC204478
VERSION CC204478.1 GI:30486459
KEYWORDS GSS.
SOURCE Gallus gallus (chicken)
ORGANISM Gallus gallus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

```

```

Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
Phasianinae; Gallus.
1 (bases 1 to 987)
Kremitzki, C., Higginbotham, J., Wylie, K., Carter, J., McPherson, J.,
Warren, W., Graves, T., Mardis, E. and Wilson, R.
Gallus gallus BAC End Reads
Unpublished (2003)
Contact: Richard K. Wilson
Genome Sequencing Center
Washington University School of Medicine
Email: submissions@wustl.edu
Insert Length: 182000 Std Error: 0.00
Seq primer: Sp6 ATTTAGTGACACTATAG
Class: BAC ends
High quality sequence start: 35
High quality sequence stop: 625.

FEATURES
source
1..987
/organism="Gallus gallus"
/mol_type="genomic DNA"
/strain="Red Jungle Fowl"
/db_xref="taxon:9031"
/clone="CH261-131C12"
/sex="female"
/cell_line="UCD001, inbred 256"
/clone_lib="CH261"
/notes="Vector: pTARBAC2.1; Site 1: EcoRI; Site 2: EcoRI;
CH261 Female Chicken library - For library and clone
ordering information: http://www.chori.org/bacpac"

ORIGIN
Query Match 81.7%; Score 18.8; DB 8; Length 987;
Best Local Similarity 90.9%; Pred. No. 7.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CATGTATTGATGGGGATAGAG 22
Db 371 CATGACTTTGATGGGGATAGAG 350

RESULT 11
BG296600/c
LOCUS
DEFINITION 602393522F1 NIH_MGC_94 Mus musculus cDNA clone IMAGE:4505236 5',
mRNA sequence.
ACCESSION BG296600
VERSION BG296600.1 GI:13059414
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 989)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgabbs-remail.nih.gov
Tissue Procurement: The Cepko Laboratory
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM10378 row: 1 column: 05
High quality sequence stop: 686.

FEATURES
source
1..989
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="IMAGE:4505236"

```

```

Db 150 GCATTTGATGGGATAGAG 131

RESULT 13
LOCUS T75599/c
DEFINITION 10377 Lambda-PRL2 Arabidopsis thaliana cDNA clone 140D23T7, mRNA
sequence.
ACCESSION T75599
VERSION T75599.1 GI:934673
KEYWORDS EST.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 365)
AUTHORS Newman,T., deBruijn,F.J., Green,P., Keegstra,K., Kende,H.,
McIntosh,L., Ohlrogge,J., Raikhel,N., Somerville,S., Thomasow,M.,
Retzel,G. and Somerville,C.
TITLE Genes galore: a summary of methods for accessing results from
large-scale partial sequencing of anonymous Arabidopsis cDNA clones
JOURNAL Plant Physiol. 106, 1241-1255 (1994)
MEDLINE 95148729
PUBMED 7846151
COMMENT On Apr 14, 1993 this sequence version replaced gi:692361.
Contact: Thomas Newman
MSU-DOE Plant Research Laboratory
Michigan State University
MSU-DOE-PRL, Michigan State University, Plant Biology Bldg., E.
Lansing, MI
Tel: 517-353-0854
Fax: 517-353-9168
Email: 22313tcn@ibm.cl.msu.edu
Seq primer: T7 dye primer.
FEATURES
Location/Qualifiers
1..365
/mol_type="mRNA"
/organism="Arabidopsis thaliana"
/ecotype="Columbia"
/db_xref="taxon:3702"
/clone="140D23T7"
/clone_lib="Lambda-PRL2"
/note="Vector: lambda Zip-Lox; Site 1: Sal; Site 2: Not;
Lambda PRL2 is a cDNA library derived from equal_
quantities of 4 pools of mRNA. The mRNA sources were 1) 7
day germinated etiolated seedlings; 2) tissue culture
grown roots; 3) staged plants half with 24 hour light
cycle, half on 16 hr light, 8 hour dark- rosettes; 4)
same plants as 3 but aerial tissue (stems, flowers and
siliques. The vector is BRL's lambda Zip-Lox. The cDNA
inserts were directionally cloned with Sal-Not arms using
oligo dt primed cDNA. "
ORIGIN
Query Match 80.0%; Score 18.4; DB 7; Length 365;
Best Local Similarity 95.0%; Pred. No. 1e+03; Indels 0; Gaps 0;
Matches 19; Conservative 0; Mismatches 1;
QY 4 GTATTTGATGGGATAGAG 23
| | | | | | | | | | | | | | | | | | | | | |
Db 81 GCATTTGATGGGATAGAG 62
| | | | | | | | | | | | | | | | | | | | | |

RESULT 14
LOCUS BP658944/c
DEFINITION BP658944 RAFL19 Arabidopsis thaliana cDNA clone RAFL19-37-J11 3',
mRNA sequence.
ACCESSION BP658944
VERSION BP658944.1 GI:49310414
KEYWORDS EST.
SOURCE Arabidopsis thaliana (thale cress)

/tissue_type="retina"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH_MGC_94"
/note="Organ: eye; Vector: pCMV-SPORT6; Site 1: NotI;
Site 2: SalI; Cloned unidirectionally; oligo-dt primed.
Average insert size 3.3 Kb. Library enriched for
full-length clones and constructed by Life Technologies.
Note: this is a NIH_MGC Library."
ORIGIN
Query Match 81.7%; Score 18.8; DB 4; Length 989;
Best Local Similarity 90.9%; Pred. No. 7.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 CATGTATTTGATGGGATAGAG 22
| | | | | | | | | | | | | | | | | | | | | |
Db 230 CATGTCTTTGATGGGATAGAG 209
| | | | | | | | | | | | | | | | | | | | | |

RESULT 12
LOCUS BH854982/c
DEFINITION SALK_087111.44.20.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_087111.44.20.x, genomic
survey sequence.
ACCESSION BH854982
VERSION BH854982.1 GI:21704572
KEYWORDS GSS.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 323)
AUTHORS Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
Gadriab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
Shinn,P., Zimmerman,J. and Ecker,J.R.
TITLE A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
JOURNAL Unpublished (2001)
COMMENT Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGNAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA. This sequence lies within 300 bases of the 3' end of
At3g02030 and an annotated exon of At3g02040.
Class: TDNA tagged.
FEATURES
Location/Qualifiers
1..323
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/ecotype="Col-0"
/db_xref="taxon:3702"
/clone="SALK_087111.44.20.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"
ORIGIN
Query Match 80.0%; Score 18.4; DB 8; Length 323;
Best Local Similarity 95.0%; Pred. No. 1e+03; Indels 1; Gaps 0;
Matches 19; Conservative 0; Mismatches 1;
QY 4 GTATTTGATGGGATAGAG 23
| | | | | | | | | | | | | | | | | | | | | |

```

ORGANISM

Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsids.
 1 (bases 1 to 372)
 Seki,M., Narusaka,M., Kamiya,A., Ishida,J., Satou,M., Sakurai,T.,
 Nakajima,M., Enju,A., Akiyama,K., Ono,Y., Muramatsu,M.,
 Hayashizaki,Y., Kawai,J., Carninci,P., Itoh,M., Ishii,Y.,
 Arakawa,T., Shibata,K., Shinagawa,A. and Shinozaki,K.
 Functional annotation of a full-length Arabidopsis cDNA collection
 Science 296 (5565), 141-145 (2002)
 21932900
 11910074

COMMENT

Contact: Motoaki Seki
 Plant Functional Genomics Research Group
 RIKEN Genomic Sciences Center
 3-1-1 Koyadai, Tsukuba, Ibaraki 305-0074, Japan
 Tel: 81-298-36-4359
 Fax: 81-298-36-9060

Email: msek@rtc.riken.go.jp

reversed clone; Please visit our web site
 (http://pfweb.gsc.riken.go.jp/) for further details.

FEATURES

source

1..372
 /organism="Arabidopsis thaliana"
 /mol_type="mRNA"
 /db_xref="taxon:3702"
 /clone="RAFL19-37-J11"
 /tissue_type="mixture of silique and flower"
 /lab_host="DH10B"
 /clone_lib="RAFL19"
 /note="Site_1: BamHI; Site_2: SalI; Subtraction Library"

ORIGIN

Query Match 80.0%; Score 18.4; DB 5; Length 372;
 Best Local Similarity 95.0%; Pred. No. 1e+03;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 GTATTGTGGGGATAGAGG 23
 | ||||| ||||| ||||| |||||

DB 303 GCATTGTGGGGATAGAGG 284
 | ||||| ||||| ||||| |||||

RESULT 15

AV814465/c

LOCUS
 DEFINITION AV814465 RAFL9 Arabidopsis thaliana cDNA clone RAFL09-83-B15 3',
 mRNA sequence.

ACCESSION

AV814465

VERSION

AV814465.1 GI:19856257

KEYWORDS

SOURCE

ORGANISM

Arabidopsis thaliana (thale cress)

Arabidopsis thaliana

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsids.

1 (bases 1 to 400)

Seki,M., Narusaka,M., Ishida,J., Kamiya,A., Satou,M., Nakajima,M.,
 Ono,Y., Sakurai,T., Carninci,P., Kawai,J., Itoh,M., Ishii,Y.,
 Arakawa,T., Shibata,K., Shinagawa,A., Muramatsu,M., Hayashizaki,Y.
 and Shinozaki,K.

Large scale analysis of Arabidopsis full-length cDNA (2002b)

Unpublished (2002)

COMMENT

Contact: Motoaki Seki
 Plant Functional Genomics Research Group
 RIKEN Genomic Sciences Center
 3-1-1 Koyadai, Tsukuba, Ibaraki 305-0074, Japan
 Tel: 81-298-36-4359
 Fax: 81-298-36-9060

Email: msek@rtc.riken.go.jp
 An Arabidopsis full-length cDNA library was constructed essentially
 as reported previously (Seki et al., 1998). cDNA cleaved with BamHI
 and XhoI was ligated to modified Lambda FLC-1 vector (Carninci et

al., submitted for publication) digested with BamHI and SalI. This
 clone is in a modified pBluescript vector. Please visit our web
 site (http://www.gsc.riken.go.jp/e/plant/index_e.html) for further
 details.

FEATURES

source

Location/Qualifiers
 1..400
 /organism="Arabidopsis thaliana"
 /mol_type="mRNA"
 /db_xref="taxon:3702"
 /clone="RAFL09-83-B15"
 /dev_stage="plants at various developmental stages from
 germination to mature seeds"
 /lab_host="DH10B"
 /clone_lib="RAFL19"
 /note="Site_1: BamHI; Site_2: SalI; subjected to
 dehydration (1, 2, 5, 10, 24 hr) and cold (1, 2, 5, 10, 24
 hr) treatments"

ORIGIN

Query Match 80.0%; Score 18.4; DB 1; Length 400;
 Best Local Similarity 95.0%; Pred. No. 1e+03;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 GTATTGTGGGGATAGAGG 23
 | ||||| ||||| ||||| |||||

DB 314 GCATTGTGGGGATAGAGG 295
 | ||||| ||||| ||||| |||||

Search completed: August 13, 2005, 06:44:49
 Job time : 1486.12 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 13, 2005, 01:08:33 ; Search time 725.234 Seconds
(without alignments)
1536.704 Million cell updates/sec

Title: US-10-673-854-2

Perfect score: 23
Sequence: 1 ccttttcacccaactaccactga 23

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.*
1: gb_ba.*
2: gb_htg.*
3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_sts.*
12: gb_sy.*
13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	23	100.0	68726	9 AC016681	AC016681 Homo sapi
C 2	23	100.0	100545	9 AL590379	AL590379 Human DNA
C 3	23	100.0	129357	9 AL590492	AL590492 Human DNA
C 4	21.4	93.0	59272	9 AL592490	AL592490 Human DNA
C 5	19.4	84.3	93279	9 AL807757	AL807757 Human DNA
C 6	18.8	81.7	155981	2 AC073361	AC073361 Homo sapi
C 7	18.8	81.7	157122	2 AC102699	AC102699 Mus muscu
C 8	18.8	81.7	165358	9 AL353782	AL353782 Homo sapi
C 9	18.8	81.7	166680	9 AC010132	AC010132 Homo sapi
C 10	18.8	81.7	174558	2 CR382283	CR382283 Danio rer
C 11	18.8	81.7	181241	9 AC080128	AC080128 Homo sapi
C 12	18.8	81.7	184801	2 CR354436	CR354436 Danio rer
C 13	18.8	81.7	187017	2 AC010101	AC010101 Homo sapi
C 14	18.8	81.7	192318	9 AC011092	AC011092 Homo sapi
C 15	18.8	81.7	193528	10 AL772234	AL772234 Mouse DNA
C 16	18.8	81.7	204548	5 AC145960	AC145960 Gallus ga
C 17	18.4	80.0	190996	10 AL671976	AL671976 Mouse DNA
C 18	18.4	80.0	207077	2 AC097903	AC097903 Rattus no
C 19	18.4	80.0	226926	10 AC122287	AC122287 Mus muscu

C 20	18.4	80.0	267058	2 AC134185	AC134185 Rattus no
C 21	18.2	79.1	549	5 AY323570	AY323570 Euplectes
C 22	18.2	79.1	1041	5 AY329424	AY329424 Muscicapa
C 23	18.2	79.1	1537	9 AF221594	AF221594 Homo sapi
C 24	18.2	79.1	3001	6 BD265960	BD265960 Schizophr
C 25	18.2	79.1	3001	6 AR244490	AR244490 Sequence
C 26	18.2	79.1	30436	9 AC093016	AC093016 Homo sapi
C 27	18.2	79.1	94847	2 AC091875	AC091875 Homo sapi
C 28	18.2	79.1	101138	9 AC027006	AC027006 Homo sapi
C 29	18.2	79.1	116983	8 OSJN00051	AL606613 Oryza sat
C 30	18.2	79.1	117048	2 AC109629	AC109629 Mus muscu
C 31	18.2	79.1	113952	10 AL731893	AL731893 Mouse DNA
C 32	18.2	79.1	144665	9 CNS07EF7	AL523558 Human chr
C 33	18.2	79.1	146313	8 OSJN00261	AL731617 Oryza sat
C 34	18.2	79.1	151509	9 CNS05TE8	AL358337 Human chr
C 35	18.2	79.1	166966	10 AC122015	AC122015 Mus muscu
C 36	18.2	79.1	167030	2 AC025646	AC025646 Homo sapi
C 37	18.2	79.1	174437	10 AC107781	AC107781 Mus muscu
C 38	18.2	79.1	181541	9 AL136524	AL136524 Human DNA
C 39	18.2	79.1	188755	9 AC013562	AC013562 Homo sapi
C 40	18.2	79.1	189897	10 AC138769	AC138769 Mus muscu
C 41	18.2	79.1	201508	2 AC026290	AC026290 Homo sapi
C 42	18.2	79.1	213785	2 CR788255	CR788255 Danio rer
C 43	18.2	79.1	225990	2 AC123627	AC123627 Mus muscu
C 44	18.2	79.1	241913	2 AC094048	AC094048 Rattus no
C 45	18.2	79.1	242318	10 AC097354	AC097354 Mus muscu

ALIGNMENTS

RESULT 1	AC016681/c	68726 bp	DNA	linear	PRI 30-SEP-2000
LOCUS	AC016681	Homo sapiens BAC clone RP11-62H15 from Y, complete sequence.			
DEFINITION	AC016681	Homo sapiens BAC clone RP11-62H15 from Y, complete sequence.			
ACCESSION	AC016681.2	GI:7321924			
VERSION	HTG.				
KEYWORDS	HTG.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	1 (bases 1 to 68726)				
AUTHORS	Sulston,J.E. and Waterston,R.				
TITLE	Toward a complete human genome sequence				
JOURNAL	Genome Res. 8 (11), 1097-1108 (1998)				
MEDLINE	99083792				
PUBMED	9847074				
REFERENCE	2 (bases 1 to 68726)				
AUTHORS	Joshu,C., Strommatt,C. and Wedgeworth,P.				
TITLE	The sequence of Homo sapiens BAC clone RP11-62H15				
JOURNAL	Unpublished				
REFERENCE	3 (bases 1 to 68726)				
AUTHORS	Waterston,R.H.				
TITLE	Direct Submission				
JOURNAL	Submitted (04-DEC-1999) Genome Sequencing Center, Washington				
REFERENCE	4 (bases 1 to 68726)				
AUTHORS	Waterston,R.H.				
TITLE	Direct Submission				
JOURNAL	Submitted (24-MAR-2000) Genome Sequencing Center, Washington				
REFERENCE	5 (bases 1 to 68726)				
AUTHORS	Waterston,R.H.				
TITLE	Direct Submission				
JOURNAL	Submitted (04-APR-2000) Genome Sequencing Center, Washington				
REFERENCE	6 (bases 1 to 68726)				
AUTHORS	Waterston,R.H.				
TITLE	Direct Submission				

JOURNAL

Submitted (07-APR-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
7 (bases 1 to 68726)

REFERENCE
AUTHORS

Waterston,R.

JOURNAL
TITLE

Submitted (30-SEP-2000) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Mar 24, 2000 this sequence version replaced gi:6524399.

COMMENT

----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc>
Contact: sapiens@wustl.edu
----- Summary Statistics

Center project name: H_NH0062H15

NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
clone sections once, or longer because we provide a small overlap
between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate
chemistry, or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by sequence
from more than one subclone; and the assembly was confirmed by
restriction digest.

MAPPING INFORMATION:

The position of this clone was established as part of a
collaboration between the Human Chromosome Y Mapping Project
(Tomoko Kawaguchi, Helen Skaletsky, Laura G. Brown, Steve Rosen,
and David C. Page at the Whitehead Institute for Biomedical
Research, Cambridge MA) and the Washington University Genome
Sequencing Center, St. Louis MO.

SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male
donor, as described by Osoegawa,K., Woon,P.Y., Zhao,B., Frengen,E.,
Tateno,M., Catanesi,J.J. and de Jong,P.J. (1998) An improved
approach for construction of bacterial artificial chromosome
libraries. Genomics 51:1-8. The clone may be obtained either from
Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong
and coworkers at the Roswell Park Cancer Institute
(<http://bacpac.med.buffalo.edu>)

VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-362J16; the clone sequenced
to the right is RP11-218E11. Actual start of this clone is at base
position 1 of RP11-62H15.

FEATURES

source

1. .68726
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="Genomic DNA"
/db_xref="taxon:9606"
/chromosome="Y"
/map="Y"

/clone="RP11-62H15"
/clone_lib="RPCI-11"
1. .352

repeat_region

1. .352

/rpt_family="L1"

repeat_region

357. .651

/rpt_family="Alu"

repeat_region

676. .769

/rpt_family="L1"

repeat_region

1837. .1873

/rpt_family="AT_rich"

repeat_region

2274. .2301

/rpt_family="AT_rich"

repeat_region

2542. .2572

/rpt_family="AT_rich"
3768. .3973
/rpt_family="MaLR"
4258. .4660
/rpt_family="MaLR"
5234. .5568
/rpt_family="MER1_type"
6159. .6713
/rpt_family="L1"
6718. .6943
/rpt_family="L1"
6929. .7694
/rpt_family="L1"
7709. .7887
/rpt_family=" (TA)n"
7903. .8247
/rpt_family="L1"
8264. .9031
/rpt_family="L1"
9435. .9583
/rpt_family="MIR"
9584. .9738
/rpt_family="MIR"
10428. .10482
/rpt_family="L2"
10574. .11996
/rpt_family="L1"
11997. .12024
/rpt_family="AT_rich"
12040. .13387
/rpt_family="L1"
13390. .13846
/rpt_family="L1"
13837. .14070
/rpt_family="L1"
14074. .15649
/rpt_family="L1"
15648. .16717
/rpt_family="L1"
16720. .16789
/rpt_family="A-rich"
16827. .17303
/rpt_family="L1"
17417. .17575
/rpt_family="L2"
17725. .17768
/rpt_family="AT_rich"
18047. .18132
/rpt_family="L1"
18338. .18648
/rpt_family="Alu"
19586. .19620
/rpt_family=" (CA)n"
19704. .19918
/rpt_family="MIR"
19925. .19980
/rpt_family=" (ATG)n"
20305. .20330
/rpt_family=" (CA)n"
20333. .20671
/rpt_family="MER2_type"
21471. .21557
/rpt_family="MaLR"
21661. .22655
/rpt_family="L1"
22725. .22971
/rpt_family="L1"
23441. .23462
/rpt_family="AT_rich"
23982. .24011
/rpt_family="AT_rich"
24167. .24258
/rpt_family="L1"


```

repeat_region 24399..24463 /rpt_family="AT_rich"
repeat_region 24468..24576 /rpt_family="L2"
repeat_region 24754..25021 /rpt_family="Retroviral"
repeat_region 25115..25575 /rpt_family="L2"
repeat_region 25715..26221 /rpt_family="MaLR"
repeat_region 26501..26523 /rpt_family="AT_rich"
repeat_region 26567..26925 /rpt_family="MaLR"

Query Match 100.0%; Score 23; DB 9; Length 68726;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCAACTACCACTGA 23
|||||
Db 57136 CCTTTTCATCCAACTACCACTGA 57114
|||||

```

```

RESULT 2
AL590379/c
LOCUS
DEFINITION Human DNA sequence from clone RP11-343H6 on chromosome X, complete
sequence.
ACCESSION AL590379
VERSION AL590379.7 GI:29466482
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 100545)
AUTHORS Howden, P.
TITLE Direct Submission
JOURNAL Submitted (01-APR-2003) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Apr 1, 2003 this sequence version replaced gi:289333281.
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk
-----

```

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest, except on the rare occasion of the clone being a YAC.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em., EMBL; Sw., SWISSPROT; Tr., TrEMBL; Wp., WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/ChrX>

RP11-343H6 is from the library RPCI-11.2 constructed by the group

of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>
VECTOR: pBACE3.6.

FEATURES

source

```

1..100545
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="X"
/clone="RP11-343H6"
/clone_lib="RPCI-11.2"

```

ORIGIN

```

Query Match 100.0%; Score 23; DB 9; Length 100545;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCAACTACCACTGA 23
|||||
Db 67668 CCTTTTCATCCAACTACCACTGA 67646
|||||

```

RESULT 3

```

AL590492/c
LOCUS
DEFINITION Human DNA sequence from clone RP11-88H5 on chromosome X, complete
sequence.
ACCESSION AL590492
VERSION AL590492.7 GI:15072594
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 129357)
AUTHORS Heath, P.
TITLE Direct Submission
JOURNAL Submitted (10-DEC-2001) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Aug 1, 2001 this sequence version replaced gi:13929489.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. The following
abbreviations are used to associate primary accession numbers given
in the feature table with their source databases: Em., EMBL; Sw.,
SWISSPROT; Tr., TrEMBL; Wp., WORMPEP; Information on the WORMPEP
database can be found at
http://www.sanger.ac.uk/Projects/C\_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome X, constructed by the Sanger Centre Chromosome X Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/ChrX
RP11-88H5 is from the library RPCI-11.1 constructed by the group of
Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACE3.6

```

COMMENT

IMPORTANT: This sequence is not the entire insert of clone RP11-88H5 it may be shorter because we sequence overlapping sections only once, except for a short overlap. The true left end of clone RP11-88H5 is at 1 in this sequence. The true left end of clone RP11-156J23 is at 129258 in this sequence.

FEATURES

Location/Qualifiers

source	1. .129357 /organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606" /chromosome="X" /clone="RP11-88H5" /clone_lib="RP11-11.1" 2. .1884 /note="L1PB2 repeat: matches 4280. .6155 of consensus" repeat_region	repeat_region	17251. .17506 /note="L1ME3A repeat: matches 5638. .5903 of consensus" 17546. .17932 /note="L1M4 repeat: matches 251. .627 of consensus" 18147. .18210 /note="L1M4 repeat: matches 616. .674 of consensus" 18184. .19717 /note="L1M4 repeat: matches 926. .2109 of consensus" 19782. .19970 /note="L1M4 repeat: matches 2123. .2308 of consensus" 19975. .20288 /note="AluJo repeat: matches 12. .312 of consensus" 20314. .20637 /note="L1M4 repeat: matches 2316. .2663 of consensus" 20644. .21468 /note="L1P3 repeat: matches 15. .650 of consensus" 21469. .23040 /note="L1PA13 repeat: matches 4584. .6152 of consensus" complement(23279. .23643) /note="match: GSS; Em:B79974" misc_feature
repeat_region	20314. .20637 /note="L1M4 repeat: matches 2316. .2663 of consensus" 20644. .21468 /note="L1P3 repeat: matches 15. .650 of consensus" 21469. .23040 /note="L1PA13 repeat: matches 4584. .6152 of consensus" complement(23279. .23643) /note="match: GSS; Em:B79974" misc_feature	repeat_region	23281. .23580 /note="AluJo repeat: matches 1. .291 of consensus" 23909. .24217 /note="AluSx repeat: matches 1. .309 of consensus" 24276. .24322 /note="WADL1 repeat: matches 33. .79 of consensus" 24326. .24482 /note="L1 repeat: matches 1992. .2147 of consensus" 24990. .25140 /note="L1M4 repeat: matches 2404. .2560 of consensus" 25585. .25983 /note="L1 repeat: matches 3095. .3404 of consensus" 25903. .26200 /note="AluYa5 repeat: matches 1. .296 of consensus" 26529. .26685 /note="PRAM repeat: matches 4. .152 of consensus" 26825. .26996 /note="MLT1J repeat: matches 320. .509 of consensus" 27032. .27116 /note="HERV16 repeat: matches 2. .89 of consensus" 27137. .27444 /note="L1TR16A repeat: matches 115. .426 of consensus" 28310. .28924 /note="L1M4 repeat: matches 5064. .5692 of consensus" 29682. .29969 /note="AluSx repeat: matches 1. .388 of consensus" 30022. .30136 /note="L1ME1 repeat: matches 6033. .6148 of consensus" 30157. .34776 /note="L1PA13 repeat: matches 648. .5299 of consensus" 34850. .35692 /note="L1PA13 repeat: matches -651. .215 of consensus" 35729. .36642 /note="L1 repeat: matches 3630. .4551 of consensus" 36652. .36809 /note="MLT1A1 repeat: matches 1. .160 of consensus" 36810. .38853 /note="L1P repeat: matches 3236. .5271 of consensus" 38877. .39968 /note="L1P repeat: matches 2318. .3413 of consensus" 40002. .40330 /note="L1P repeat: matches 2201. .2320 of consensus" 40118. .40327 /note="AluY repeat: matches 94. .303 of consensus" 40328. .41566 /note="L1P repeat: matches 2656. .3894 of consensus" 41571. .41865 /note="AluDb repeat: matches 1. .288 of consensus" 41890. .41983 /note="L1 repeat: matches 3856. .3955 of consensus" 44982. .45187 /note="MLT1 repeat: matches 1. .205 of consensus" 45472. .45874

```

/notes="MLTII repeat: matches 1. .402 of consensus"
46448. .46782
/notes="MER58B repeat: matches 2. .341 of consensus"
47373. .48157
/notes="LI repeat: matches 4164. .4992 of consensus"

Query Match      100.0%; Score 23; DB 9; Length 129357;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCCACTACCACTGA 23
    |||||
Db 98349 CCTTTTCATCCCACTACCACTGA 98327

RESULT 4
AL592490 59272 bp DNA linear PRI 25-OCT-2001
LOCUS Human DNA sequence from clone RP11-432J3 on chromosome 13, complete
DEFINITION
ACCESSION AL592490
VERSION AL592490.8 GI:16501199
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
TITLE 1 (bases 1 to 59272)
JOURNAL Ramsay, H.
COMMENT Direct Submission
Submitted (25-OCT-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
requests: clonerequest@sanger.ac.uk
On Oct 26, 2001 this sequence version replaced gi:16416251.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. The following
abbreviations are used to associate primary accession numbers given
in the feature table with their source databases: Em:, EMBL; Sw:,
SWISSPROT; Tr:, TrEMBL; Wp:, WORMPEP; Information on the WORMPEP
database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome 13, constructed by the Sanger Centre Chromosome 13
Mapping Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr13
RP11-432J3 is from the library RPCI-11.2 constructed by the group
of Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACE3.6
IMPORTANT: This sequence is not the entire insert of clone
RP11-432J3 It may be shorter because we sequence overlapping
sections only once, except for a short overlap.
The true left end of clone RP11-210L5 is at 57273 in this sequence.
The true right end of clone RP11-310K10 is at 2000 in this
sequence.

FEATURES             Location/Qualifiers
     source          1..59272
                     /organism="Homo sapiens"
                     /mol_type="genomic DNA"
                     /db_xref="taxon:9606"
                     /chromosome="13"
                     /clone="RP11-432J3"

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/misc_feature      /clone_lib="RPCI-11.2"
2001. .2122
/notes="Single clone region. Assembly confirmed by
restriction digest data."
misc_feature      2298. .2442
/notes="Single clone region. Assembly confirmed by
restriction digest data."
misc_feature      24343. .24487
/notes="Single clone region. Assembly confirmed by
restriction digest data."
misc_feature      28864. .28869
/notes="1327 bases of IS2 transposon (V00610) removed here.
This sequence represents the duplicated flanking sequence
of the IS2."

ORIGIN
Query Match      93.0%; Score 21.4; DB 9; Length 59272;
Best Local Similarity 95.7%; Pred. No. 9.8;
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCCACTACCACTGA 23
    |||||
Db 31913 CCTTTTCATCCCACTACCACTGA 31935

RESULT 5
AL807757/c 93279 bp DNA linear PRI 17-NOV-2002
LOCUS Human DNA sequence from clone RP11-490L4 on chromosome 9, complete
DEFINITION
ACCESSION AL807757
VERSION AL807757.9 GI:25137169
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
TITLE 1 (bases 1 to 93279)
JOURNAL Direct Submission
Submitted (16-NOV-2002) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Nov 19, 2002 this sequence version replaced gi:23895171.
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk
-----
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em:, EMBL; Sw:, SWISSPROT; Tr:, TrEMBL; Wp:, WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr9
RP11-490L4 is from the library RPCI-11.1 constructed by the group
of Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACE3.6
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such

```

as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest, except on the rare occasion of the clone being a YAC.

FEATURES

source

```

1. .93279
   Location/Qualifiers
     /organism="Homo sapiens"
     /mol_type="genomic DNA"
     /db_xref="taxon:9606"
     /chromosome="9"
     /clone="RP11-49014"
     /clone_lib="RP11-11.1"

```

ORIGIN

```

Query Match      84.3%; Score 19.4; DB 9; Length 93279;
Best Local Similarity 95.2%; Pred. NO. 90;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY 3 TTTTCATCCAACTACCACCTGA 23

Db 19692 TTTTCATCCAACTACCACCTGA 19672

RESULT 6

AC073361/c

```

LOCUS          AC073361      155981 bp      DNA      linear      HTG 08-JAN-2003
DEFINITION     Homo sapiens chromosome 3 clone RP11-160F1, WORKING DRAFT SEQUENCE,
                20 unordered pieces.

```

ACCESSION

AC073361.8 GI:20335704

VERSION HTG; HTGS_PHASE1; HTGS_DRAFT.

KEYWORDS

Homo sapiens (human)

SOURCE

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 155981)

Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C., Alsbrooks,S.L., Amaralunge,H.C., Are,J.R., Ayete,M., Banks,T., Barbara,J., Benton,J., Bimage,K., Blankenburg,K., Bonnin,D., Bouck,J., Bowie,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C., Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Z., Chowdhry,I., Christopoulos,C., Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinh,H.H., Douthwaite,K.J., Draper,H., Dugan-Rocha,S., Durbin,K.J., Earnhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T., Garza,N., Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S., Hamilton,K., Harris,C., Harris,K., Hart,M., Havlak,P., Hawes,A., Hernandez,J., Hernandez,O., Hodgson,A., Hogue,M., Holloway,C., Hollins,B., Homs,F., Howard,S., Huber,J., Hulyk,S., Hume,J., Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudah,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvah,J., Kovar,C., Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C., Lewis,L., Li,J., Li,Z., Lichtarge,O., Lieu,C., Liu,J., Liu,W., Louseged,H., Lozado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R., Ma,J., Maheshwari,M., Mapa,P., Martin,R., Martindale,A., Martinez,E., Massey,E., Mawhiney,E., McLeod,M.P., Meador,M., Mei,G., Metzker,M., Miner,G., Miner,Z., Mitchell,T., Mohabbat,K., Morgan,M., Morris,S., Moser,M., Neal,D., Newton,J., Newton,S., Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokoko,S., Ogih,M., Okuwonu,G., Oragune,N., Oviedo,R., Pace,A., Payton,B., Peery,J., Perez,L., Peters,L., Pickens,R., Primus,E., Pu,L., Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojibokan,I., Rolfe,M., Ruiz,S., Savary,G., Scherer,S., Scott,G., Shen,H., Shooshkar,N., Sisson,I., Sodergren,E., Sonaik,T., Sparks,A., Stanley,H., Stone,H., Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Tamerisa,K., Tang,H., Tansley,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S., Usmani,K., Vasquez,L., Vera,V., Villalon,D., Vinson,R., Wang,Q., Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S.,

Williams,G., Williamson,A., Wlarczyk,R., Wooden,S., Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D., Weinstock,G. and Gibbs,R.
 Unpublished
 2 (bases 1 to 155981)
 Worley,K.C.
 Direct Submission
 Submitted (15-JUN-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 3 (bases 1 to 155981)
 Worley,K.C.
 Direct Submission
 Submitted (08-JAN-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 On Apr 28, 2002 this sequence version replaced gi:15887171.

COMMENT

```

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Drafting Center Code: BCM
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HBQ
Center clone name: RP11-160F1
----- Summary Statistics
Sequencing vector: M13;
Chemistry: Dye-Primer Bodipy: 40% of reads
Chemistry: Dye-terminator Big Dye: 60% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 141831 bases at least Q40
Consensus quality: 146281 bases at least Q30
Consensus quality: 148168 bases at least Q20
Estimated insert size: 155230; sum-of-contigs estimation
Quality coverage: 4x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: Estimated insert size may differ from sequence length
  (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
  consists of 20 contigs. The true order of the pieces
  is not known and their order in this sequence record is
  arbitrary. Gaps between the contigs are represented as
  runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
  as soon as it is available and the accession number will
  be preserved.

```

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* 1 2542: contig of 2542 bp in length
* 2543 2642: gap of unknown length
* 2643 4841: contig of 2199 bp in length
* 4842 4941: gap of unknown length
* 4942 7369: contig of 2428 bp in length
* 7370 7469: gap of unknown length
* 7470 11527: contig of 4058 bp in length
* 11528 11627: gap of unknown length
* 11628 16423: contig of 4796 bp in length
* 16424 16523: gap of unknown length
* 16524 20209: contig of 3686 bp in length
* 20210 20309: gap of unknown length
* 20310 24291: contig of 3982 bp in length
* 24292 24391: gap of unknown length
* 24392 27954: contig of 3562 bp in length
* 27954 28053: gap of unknown length
* 28054 32347: contig of 4194 bp in length
* 32348 37027: contig of 4680 bp in length
* 37028 37127: gap of unknown length
* 37128 41991: contig of 4864 bp in length
* 41992 42091: gap of unknown length
* 42092 47186: contig of 5095 bp in length
* 47187 47286: gap of unknown length
* 47287 53916: contig of 6630 bp in length

```

```

* 53917 54016: gap of unknown length
* 54017 59973: contig of 5957 bp in length
* 59974 60073: gap of unknown length
* 60074 67148: contig of 7075 bp in length
* 67149 67248: gap of unknown length
* 67249 77842: contig of 10594 bp in length
* 77843 77942: gap of unknown length
* 77943 91209: contig of 13267 bp in length
* 91210 91309: gap of unknown length
* 91310 104946: contig of 13637 bp in length
* 104947 105046: gap of unknown length
* 105047 126326: contig of 21280 bp in length
* 126327 126426: gap of unknown length
* 126427 155981: contig of 29555 bp in length.
FEATURES
    source
        1..155981
            /organism="Homo sapiens"
            /mol_type="genomic DNA"
            /db_xref="taxon:9606"
            /chromosome="3"
            /clone="RP11-160F1"
ORIGIN
Query Match      81.7%  Score 18.8;  DB 2;  Length 155981;
Best Local Similarity 90.9%;  Pred. No. 1.7e+02;
Matches 20;  Conservative 0;  Mismatches 2;  Indels 0;  Gaps 0;

QY  2  CTTTTCATCAACTACCACTGA 23
      |||||
Db  143609 CGTTTCATCCAGCTACCACTGA 143588

RESULT 7
AC102699/c
LOCUS      AC102699          157122 bp    DNA        linear      HTG 03-MAR-2003
DEFINITION Mus musculus clone RP24-268F1, WORKING DRAFT SEQUENCE, 6 unordered
            pieces.
ACCESSION  AC102699
VERSION    AC102699.3  GI:28631330
KEYWORDS   HTG; HTGS PHASE1; HTGS DRAFT; HTGS_FULLTOP.
SOURCE     Mus musculus
            (house mouse)
ORGANISM   Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE  1  (bases 1 to 157122)
            Birren,B., Nusbaum,C. and Lander,E.
            Unpublished
            Mus musculus, clone RP24-268F1
2  (bases 1 to 157122)
            Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
            Anderson,S., Barna,N., Bastien,V., Boguslavskiy,L., Boukhgalter,B.,
            Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,
            Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A.,
            Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
            Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S.,
            Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-pierre,N.,
            Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
            Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K.,
            Lamazares,R., Landers,T., Lehotsky,J., Levine,R., Liu,G.,
            MacLean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C.,
            McCarthy,M., McEwan,P., McKernan,K., McPheeters,R., Meldrim,J.,
            Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,
            Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D.,
            Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
            Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
            Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schuback,R.,
            Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
            Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
            Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
            Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
            Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
            Direct Submission
            Submitted (23-NOV-2001) Whitehead Institute/MIT Center for Genome
            Research

```

REFERENCE
AUTHORS

Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 157122)
Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N.,
Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T.,
Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y.,
Collymore,A., Cook,A., Cooke,P., Corum,B., DeArellano,K., Faro,S.,
Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Gardyna,S.,
Ferreira,P., FitzGerald,M., Gage,D., Galagan,J., Gardyna,S.,
Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B.,
Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R.,
Lindblad-Toh,K., Liu,G., Lui,A., Mabbitt,R., MacLean,C.,
Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M.,
Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J.,
Nguyen,C., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P.,
O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Schauer,S., Schuback,R., Seaman,S., Severy,P., Smith,C.,
Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M.,
Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M.,
Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X.,
Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (03-MAR-2003) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Mar 3, 2003 this sequence version replaced gi:22381689.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

TITLE

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR

JOURNAL

Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu

COMMENT

----- Project Information
Center project name: L19308
Center clone name: 268_F1

SUMMARY

----- Summary Statistics
Sequencing vector: plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 156083 bases at least Q40
Consensus quality: 156406 bases at least Q30
Consensus quality: 156521 bases at least Q20
Insert size: 151000; agarose-fp
Insert size: 156622; sum-of-contigs
Quality coverage: 10.5 in Q20 bases; agarose-fp
Quality coverage: 10.1 in Q20 bases; sum-of-contigs

NOTE

* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

FEATURES

1 51217: contig of 51217 bp in length
* 51218 51317: gap of 100 bp
* 51318 52366: contig of 1049 bp in length
* 52367 52466: gap of 100 bp
* 52467 114294: contig of 61828 bp in length
* 114295 114394: gap of 100 bp
* 114395 121686: contig of 7292 bp in length
* 121687 121786: gap of 100 bp
* 121787 153936: contig of 32150 bp in length
* 153937 154036: gap of 100 bp
* 154037 157122: contig of 3086 bp in length.

Location/Qualifiers

1..157122
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"

TITLE

Center: Whitehead Institute/MIT Center for Genome
Center code: WIBR

JOURNAL

Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu

COMMENT

----- Project Information
Center project name: L19308
Center clone name: 268_F1

SUMMARY

----- Summary Statistics
Sequencing vector: plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 156083 bases at least Q40
Consensus quality: 156406 bases at least Q30
Consensus quality: 156521 bases at least Q20
Insert size: 151000; agarose-fp
Insert size: 156622; sum-of-contigs
Quality coverage: 10.5 in Q20 bases; agarose-fp
Quality coverage: 10.1 in Q20 bases; sum-of-contigs

NOTE

* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

FEATURES

1 51217: contig of 51217 bp in length
* 51218 51317: gap of 100 bp
* 51318 52366: contig of 1049 bp in length
* 52367 52466: gap of 100 bp
* 52467 114294: contig of 61828 bp in length
* 114295 114394: gap of 100 bp
* 114395 121686: contig of 7292 bp in length
* 121687 121786: gap of 100 bp
* 121787 153936: contig of 32150 bp in length
* 153937 154036: gap of 100 bp
* 154037 157122: contig of 3086 bp in length.

Location/Qualifiers

1..157122
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"

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/misc_feature      /clone="RP24-268P1"
                  /clone_lib="RPC1-24 Male Mouse BAC"
                  1. .51217
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clone end:SP6
vector_side:left"
misc_feature      51318..52366
                  /note="assembly_fragment"
52467..114294
misc_feature      /note="assembly_fragment"
114395..121686
misc_feature      /note="assembly_fragment"
121787..153936
misc_feature      /note="assembly_fragment"
154037..157122
misc_feature      /note="assembly_fragment
clone end:T7
vector_side:right"

ORIGIN
Query Match      81.7%; Score 18.8; DB 2; Length 157122;
Best Local Similarity 90.9%; Pred. No. 1.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      2      CTTTTCATCCCACTACCACTGA 23
|||||
Db      57866 CTTTTCATCCCACTAGAACTGA 57845

RESULT 8
AL353782/c
LOCUS      AL353782
DEFINITION      Human DNA sequence from clone RP11-207C11 on chromosome 9, complete
sequence.
ACCESSION      AL353782
VERSION      AL353782.23 GI:18476550
KEYWORDS      HTG.
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Sehra,H.
Direct Submission
Submitted (30-JAN-2002) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Feb 1, 2002 this sequence version replaced gi:17127880.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. The following
abbreviations are used to associate primary accession numbers given
in the feature table with their source databases: Em:, EMBL; Swi,
SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information on the WORMPEP
database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
Chromosome 9, constructed by the Sanger Centre
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr9
RP11-207C11 is from the library RPC1-11.1 constructed by the group
of Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm

```

```

VECTOR: pBAC3.6
IMPORTANT: This sequence is not the entire insert of clone
RP11-207C11 it may be shorter because we sequence overlapping
sections only once, except for a short overlap.
The true right end of clone RP11-207C11 is at 165358 in this
sequence. The true right end of clone RP11-349P17 is at 2000 in
this sequence.
FEATURES             Location/Qualifiers
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                     /mol_type="genomic DNA"
                     /db_xref="taxon:9606"
                     /chromosome="9"
                     /clone="RP11-207C11"
                     /clone_lib="RPC1-11.1"
                     /clone_size:165358
     misc_feature     /note="Single clone region. Assembly confirmed by
restriction digest data."
                     114473..114543
     misc_feature     /note="Single clone region. Sequence from reads from a
short insert library derived from a single pUC clone.
Restriction digest data confirm the assembly."
ORIGIN
Query Match      81.7%; Score 18.8; DB 9; Length 165358;
Best Local Similarity 90.9%; Pred. No. 1.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      2      CTTTTCATCCCACTACCACTGA 23
|||||
Db      129391 CTTTTCATCCCACTCCCACTGA 129370

RESULT 9
AC010132/c
LOCUS      AC010132
DEFINITION      Homo sapiens BAC clone RP11-111K18 from 7p11.2-p2, complete
sequence.
ACCESSION      AC010132
VERSION      AC010132.5 GI:10440742
KEYWORDS      HTG.
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Suleston,J.E. and Waterston,R.
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
99063792
MEDLINE      9847074
REFERENCE     2 (bases 1 to 166680)
AUTHORS      Andrews,S., Wohlmann,P. and Scherger,E.
TITLE        The sequence of Homo sapiens BAC clone RP11-111K18
JOURNAL      Unpublished
REFERENCE     3 (bases 1 to 166680)
AUTHORS      Waterston,R.H.
TITLE        Direct Submission
JOURNAL      Submitted (13-SEP-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE     4 (bases 1 to 166680)
AUTHORS      Waterston,R.
TITLE        Direct Submission
JOURNAL      Submitted (30-SEP-2000) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Sep 30, 2000 this sequence version replaced gi:8954217.
COMMENT      ----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@watson.wustl.edu
----- Summary Statistics

```

Center project name: H_NH0111K18

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

The sequence of this clone was established as part of a mapping and sequencing collaboration between the NHGRI Chromosome 7 Mapping Project (Eric D. Green, Director), John D. McPherson in the Department of Genetics (Washington University), and the Washington University Genome Sequencing Center. For additional information about the map position of this sequence, see <http://www.nhgri.nih.gov/DIR/GRB/CHR7>, send <mailto:egreen@nhgri.nih.gov>, or see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RPC1-11 human BAC library was made from the blood of one male donor, as described by Onogawa, K., Woon, P.Y., Zhao, B., Frengen, B., Tateno, M., Catanese, J.J., and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. *Genomics* 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBACes.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is GS1-308H5, 200 bp overlap. Actual start of this clone is at base position 156485 of GS1-308H5; actual end is at base position 166680 of Rp11-111K18.

There are polymorphic base pair differences in the overlap between the clone Rp11-111K18 and GS1-308H5.

FEATURES

source	1. 166680	/organism="Homo sapiens"	/rpt_family="Alu"
		/mol_type="Genomic DNA"	/rpt_family="ERV1"
		/db_xref="taxon:9606"	4260. 4865
		/chromosome="7"	/rpt_family="ERV1"
		/map="7p11.2-p2"	4902. 4994
		/clone="Rp11-111K18"	/rpt_family="ERV1"
		/clone_lib="RPC1-11"	6077. 6610
repeat_region	2. 397	/rpt_family="ERV1"	/rpt_family="ERV1"
		1478. 1785	7185. 7327
repeat_region		/rpt_family="Alu"	/rpt_family="ERV1"
		1807. 1913	7327. 7539
repeat_region		/rpt_family="L1"	/rpt_family="ERV1"
		2138. 2508	7579. 8074
repeat_region		/rpt_family="ERV1"	/rpt_family="ERV1"
		2532. 2817	8169. 9081
repeat_region		/rpt_family="MaLR"	/rpt_family="ERV1"
		2846. 2911	9082. 9362
repeat_region		/rpt_family="ERV1"	/rpt_family="Alu"
		2912. 3043	9363. 9633
repeat_region		/rpt_family="Alu"	/rpt_family="ERV1"
		3044. 3115	9634. 9943
repeat_region		/rpt_family="ERV1"	/rpt_family="Alu"
		3087. 3598	9944. 10150
repeat_region		/rpt_family="ERV1"	/rpt_family="ERV1"
		3600. 3780	10151. 10567
repeat_region		/rpt_family="ERV1"	/rpt_family="MaLR"
		3781. 4090	10568. 12227
			/rpt_family="ERV1"
			12316. 12514
			/rpt_family="ERV1"
			12532. 12582
			/rpt_family="ERV1"
			13068. 13614
			/rpt_family="ERV1"
			15167. 15445
			/rpt_family="Alu"
			16828. 17212
			/rpt_family="MaLR"
			17548. 17857
			/rpt_family="Alu"
			17897. 18030
			/rpt_family="ERV1"
			18365. 18652
			/rpt_family="Alu"
			19118. 19746
			/rpt_family="ERV1"
			19750. 19814
			/rpt_family="ERV1"
			19852. 20169
			/rpt_family="ERV1"
			20224. 20412
			/rpt_family="ERV1"
			20540. 20840
			/rpt_family="Alu"
			20879. 20997
			/rpt_family="MIR"
			21643. 21848
			/rpt_family="MER1_type"
			22158. 22177
			/rpt_family="MIR"
			22178. 22488
			/rpt_family="ERV1"
			22489. 22777
			/rpt_family="Alu"
			22778. 22919
			/rpt_family="ERV1"
			22920. 22979
			/rpt_family="MIR"
			23031. 23067
			/rpt_family="L2"
			23518. 23928
			/rpt_family="MaLR"

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repeat_region 24091..25047
/rpt_family="L2"
repeat_region 25091..25312
/rpt_family="L2"
repeat_region 25375..25549
/rpt_family="L2"
repeat_region 25663..25817
/rpt_family="MER1_type"
repeat_region 27098..27212
/rpt_family="L2"
repeat_region 27657..27925
/rpt_family="MALR"
repeat_region 28020..28061
/rpt_family="L2"
repeat_region 28527..28780
/rpt_family="Alu"

Query Match 81.7%; Score 18.8; DB 9; Length 166680;
Best Local Similarity 90.9%; Pred. No. 1.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 CTTTTCATCCAACTACCACTGA 23
| ||||| ||||| ||||| |||||
Db 71916 CATTTCATCCAACTACCACTG 71895

RESULT 10
CR382283/c
LOCUS
DEFINITION
Danio rerio clone DKEYP-32G11, ** SEQUENCING IN PROGRESS ***, 9
unordered pieces.
ACCESSION
CR382283
VERSION
CR382283.2 GI:46019446
KEYWORDS
HTG; HTGS_PHASE1.
SOURCE
Danio rerio (zebrafish)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.
Sims,S.
Direct Submission
Submitted (29-MAR-2004) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
zf1sh-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Apr 1, 2004 this sequence version replaced gi:45774318.
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: zf1sh-help@sanger.ac.uk
----- Project Information
Center project name: zkg32G11
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Chemistry: Dye-terminator; 100% of reads
Consensus quality: 171524 bases at least Q40
Consensus quality: 172036 bases at least Q30
Consensus quality: 172486 bases at least Q20
Insert size: 173758; sum-of-contigs
Insert size: 172615; 6.9% error; agarose-fp
Quality coverage: 7.43x in Q20 bases; sum-of-contigs Quality
coverage: 7.76x in Q20 bases; agarose-fp
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 9 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 8310: contig of 8310 bp in length

repeat_region 8410: gap of 100 bp
* 8311 8410: gap of 100 bp
* 8411 52211: contig of 43801 bp in length
* 52212 52311: gap of 100 bp
* 52312 71637: contig of 19326 bp in length
* 71638 71737: gap of 100 bp
* 71738 79714: contig of 7977 bp in length
* 79715 79814: gap of 100 bp
* 79815 107925: contig of 28111 bp in length
* 107926 108025: gap of 100 bp
* 108026 139165: contig of 31140 bp in length
* 139166 139265: gap of 100 bp
* 139266 142778: contig of 3513 bp in length
* 142779 142878: gap of 100 bp
* 142879 158385: contig of 15507 bp in length
* 158386 158486: gap of 100 bp
* 158487 174558: contig of 16073 bp in length.
* 158488

FEATURES
Location/Qualifiers
source
1..174558
/organism="Danio rerio"
/mol_type="genomic DNA"
/db_xref="taxon:7955"
/clone="DKEYP-32G11"
/clone_lib="DanioKeyPilot"
1..8310
/note="assembly fragment:00142
fragment_chain:1"
8411..52211
/note="assembly fragment:01851
fragment_chain:1"
52312..71637
/note="assembly fragment:00673
fragment_chain:1"
71738..79714
/note="assembly fragment:00067
fragment_chain:1"
79815..107925
/note="assembly fragment:01363
fragment_chain:1"
108026..139165
/note="assembly fragment:00960
fragment_chain:1"
139266..142778
/note="assembly fragment:00035.0"
142879..158385
/note="assembly fragment:00253"
158486..174558
/note="assembly fragment:00432"

ORIGIN
Query Match 81.7%; Score 18.8; DB 2; Length 174558;
Best Local Similarity 90.9%; Pred. No. 1.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CTTTTCATCCAACTACCACTG 22
||||| ||||| ||||| |||||
Db 78871 CCTTTATCCAGCTACCACTG 78850

RESULT 11
AC080128
LOCUS
DEFINITION
Homo sapiens 3 BAC RP11-404G16 (Roswell Park Cancer Institute Human
BAC Library) complete sequence.
ACCESSION
AC080128
VERSION
AC080128.21 GI:21591797
KEYWORDS
HTG.
SOURCE
Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 181241)
Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,
Alsbrooks,S.L., Amarantunge,H.C., Are,J.R., Ayele,M., Banks,T.,

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repeat_region complement(14585..14886)
repeat_region /rpt_family="LIME2"
repeat_region complement(15313..15378)
repeat_region /rpt_family="LIMD2"
repeat_region 15379..15691
repeat_region /rpt_family="AluSq"
repeat_region complement(15692..15955)
repeat_region /rpt_family="LIMD2"
repeat_region complement(15956..16258)
repeat_region /rpt_family="AluSx"
repeat_region complement(16259..16359)
repeat_region /rpt_family="LIMD2"
repeat_region 16442..16603
repeat_region /rpt_family="LIME2"
repeat_region complement(16640..16811)
repeat_region /rpt_family="LIMD1"
repeat_region complement(16812..17088)
repeat_region /rpt_family="LIPa7"
repeat_region complement(17089..17745)
repeat_region /rpt_family="LIMD1"
repeat_region complement(17752..17990)
repeat_region /rpt_family="LIME"
repeat_region complement(18007..18439)
repeat_region /rpt_family="LIMEc"
repeat_region 18446..18493
repeat_region /rpt_family="AT rich"
repeat_region complement(18532..19199)
repeat_region /rpt_family="LIMEc"
repeat_region complement(19437..19934)
repeat_region /rpt_family="LIMCc"
repeat_region 19974..20595
repeat_region /rpt_family="LIME2"
repeat_region 20632..20968
repeat_region /rpt_family="LIME2"
repeat_region 20875..20981
repeat_region /standard_name="D3S2333"
repeat_region 20969..21275
repeat_region /rpt_family="AluSx"
repeat_region 21276..21288
repeat_region /rpt_family="LIME2"
repeat_region complement(21921..22221)

Query Match 81.7%; Score 18.8; DB 9; Length 181241;
Best Local Similarity 90.9%; Pred. No. 1.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 CTTTTCATCCAACTACCACCTGA 23
Db 106165 CGTTTCATCCAGCTACCACCTGA 106186

RESULT 12
CR354436 184801 bp DNA linear HTG 10-OCT-2004
LOCUS Danio rerio clone DKEY-269F18, WORKING DRAFT SEQUENCE, 4 unordered
DEFINITION Pieces.
ACCESSION CR354436
VERSION CR354436.8 GI:54021885
KEYWORDS HTG; HTGS_PHASE1; HTGS_ACTIVEFIN; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.
REFERENCE 1 (bases 1 to 184801)
AUTHORS Almeida, J.
TITLE Direct Submission
JOURNAL Submitted (08-OCT-2004) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
zf1sh-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
COMMENT On Oct 10, 2004 this sequence version replaced gi:53142909.
----- Genome Center
Center: Wellcome Trust Sanger Institute

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```

Center code: SC
Web site: http://www.sanger.ac.uk
Contact: zf1sh-help@sanger.ac.uk
----- Project Information
Center project name: zk269F18
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Chemistry: Dye-terminator; 100% of reads
Consensus quality: 184321 bases at least Q40
Consensus quality: 184412 bases at least Q30
Consensus quality: 184448 bases at least Q20
Insert size: 184501; sum-of-contigs
Quality coverage: 9.52x in Q20 bases; sum-of-contigs Quality
Coverage: 10.35x in Q20 bases; agarose-fp
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 4 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 154052: contig of 154052 bp in length
* 154053 154152: gap of 100 bp
* 154153 161873: contig of 7721 bp in length
* 161874 161973: gap of 100 bp
* 161974 181767: contig of 19794 bp in length
* 181768 181867: gap of 100 bp
* 181868 184801: contig of 2934 bp in length.
* 181868 184801: Location/Qualifiers
* 1..184801
* /organism="Danio rerio"
* /mol_type="genomic DNA"
* /db_xref="taxon:7955"
* /clone="DKEY-269F18"
* /clone_lib="DanioKey"
* 1..154052
* /note="assembly fragment:02802"
* fragment_chain:1
* 154153..161873
* /note="assembly fragment:01702"
* fragment_chain:1
* 161974..181767
* /note="assembly fragment:01449"
* fragment_chain:1
* 181868..184801
* /note="assembly fragment:02662"

ORIGIN

Query Match 81.7%; Score 18.8; DB 2; Length 184801;
Best Local Similarity 90.9%; Pred. No. 1.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CTTTTCATCCAACTACCACCTG 22
Db 31456 CGTTTCATCCAGCTACCACCTG 31477

RESULT 13
AC010101/c 187017 bp DNA linear HTG 08-AUG-2000
LOCUS Homo sapiens chromosome 7 clone RP11-447C21, WORKING DRAFT
DEFINITION SEQUENCE, 8 unordered pieces.
ACCESSION AC010101
VERSION AC010101.8 GI:9739351
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 187017)

```

AUTHORS	Waterston, R.H.
TITLE	The sequence of Homo sapiens clone
JOURNAL	Unpublished
REFERENCE	2 (bases 1 to 187017)
AUTHORS	Waterston, R.H.
TITLE	Direct Submission
JOURNAL	Submitted (11-SEP-1999) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
COMMENT	On Aug 8, 2000 this sequence version replaced qi:9653201.

```

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H NH0447C21
----- Summary Statistics -----
Sequencing vector: M13; 63%
Chemistry: Dye-primer ET; 48% of reads
Assembly: Dye-terminator Big Dye; 52% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 181338 bases at least Q40
Consensus quality: 183284 bases at least Q30
Consensus quality: 183284 bases at least Q20
Insert size: 190000; agarose-fp
Insert size: 186317; sum-of-contigs
Quality coverage: 6.29 in Q20 bases; agarose-fp
Quality coverage: 6.62 in Q20 bases; sum-of-contigs

```

* NOTE. This is a 'working draft' sequence. It currently
* consists of 8 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence,
* as soon as it is available and the accession number will
* be preserved.

*	1	4028:	contig of 4028 bp in length
*	4029	4128:	gap of unknown length
*	4129	7967:	contig of 3839 bp in length
*	7968	8067:	gap of unknown length
*	8068	16351:	contig of 8284 bp in length
*	16352	16451:	gap of unknown length
*	16452	26437:	contig of 9986 bp in length
*	26438	26537:	gap of unknown length
*	26538	35888:	contig of 9351 bp in length
*	35889	35988:	gap of unknown length
*	35989	56349:	contig of 20361 bp in length
*	56350	56449:	gap of unknown length
*	56350	85530:	contig of 29081 bp in length
*	85531	85630:	gap of unknown length
*	85631	187017:	contig of 101387 bp in length

FEATURES
SOURCE

```

location/Quarrels
1. .187017
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="7"
/clone="RP11-447C21"

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misc_feature      1. .4028
                  /notes="assembly_name:Contig26"
misc_feature      4129. .7957
                  /notes="assembly_name:Contig27"
misc_feature      8068. .16351
                  /note="assembly_name:Contig28"
misc_feature      16452. .28437
                  /notes="assembly_name:Contig29
                  clone_end:T7
misc_feature      28538. .35888
                  vector_side:left"
misc_feature      /notes="assembly_name:Contig30"
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<hr/>					
misc_feature	35989..56349				
	/note="assembly_name:Contig31"				
misc_feature	56450..85530				
	/note="assembly_name:Contig32				
	clone_end:SP6				
	vector_side:right"				
misc_feature	85631..187017				
	/note="assembly_name:Contig33"				
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ORIGIN					
Query Match	81.7%	Score 18.8;	DB 2;	Length 187017;	
Best Local Similarity	90.9%	Pred. No. 1.7e+02;			
Matches 20;	Conservative 0;	Mismatches 2;	Indels 0;	Gaps 0;	
<hr/>					
Qy	2 CTTTTCATCCAACTACCACGTGA 23				
Dd	123519 CATTTTCATCCAACTACCACAGA 123498				
<hr/>					
RESULT 14					
ACOL1092/c					
LOCUS	Homo sapiens, clone RP11-1H15, complete sequence.	192318 bp	DNA linear	PRI 12-JAN-2002	
DEFINITION	ACOL1092				
ACCESSION	ACOL1092.4 GI:18034740				
VERSION	HTG.				
KEYWORDS	Homo sapiens (human)				
SOURCE	Homo sapiens				
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
REFERENCE	1 (bases 1 to 192318)				
AUTHORS	Birren,B., Linton,L., Nussbaum,C. and Lander,E.				
TITLE	Homo sapiens, clone RP11-1H15				
JOURNAL	Unpublished				
REFERENCE	2 (bases 1 to 192318)				
AUTHORS	Birren,B., Linton,L., Nussbaum,C., Lander,E., Allen,N., Anderson,M., Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Boukhgalter,B., Brown,A., Castile,A., Colangelo,M., Collins,S., Collymore,A., Cooke,P., DeArellano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M., Ferreira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D., Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L., Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J., Lehoczyk,J., Lieu,C., Locke,K., MacDonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J., Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P., Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P., Strange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J., Testfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X., Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.				
TITLE	Direct Submission				
JOURNAL	Submitted (01-OCT-1999) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA				
REFERENCE	3 (bases 1 to 192318)				
AUTHORS	Birren,B., Linton,L., Nussbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Boukhgalter,B., Brown,A., Camarata,J., Campoianno,A., Chang,J., Charazo,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A., Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S., Fero,S., Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S., Glinde,S., Gord,S., Guyette,M., Graham,L., Grand-Pierre,N., Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K., Lamarez,R., Landers,T., Lehoczyk,J., Levine,R., Liu,G., MacLean,C., MacDonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K., MCPheeters,R., Meldrim,J., Menues,L., Mihova,T., Mienga,V., Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,C., Schuback,R., Seaman,S., Severy,P., Spencer,B., Strange-Thomann,N., Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Testfaye,S., Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,				

Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G.,
 Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
 Direct Submission
 Submitted (12-JAN-2002) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Jan 2, 2002 this sequence version replaced gi:12718900.
 All repeats were identified using RepeatMasker:
 Smit, A. F. A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html
 ----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu
 ----- Project Information
 Center project name: L1460
 Center clone name: 1_H_15

FEATURES

Location/Qualifiers

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Search completed: August 13, 2005, 05:04:15
Job time : 732.234 secs

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Best Local Similarity 90.9%; Pred. No. 1.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 CTTTTCATCCAACTACCACTGA 23
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Db 72240 CATTTCATCCAACTACCACTGA 72219

RESULT 15
AL772234/c
LOCUS
DEFINITION AL772234 193528 bp DNA linear ROD 01-OCT-2002
Mouse DNA sequence from clone RP23-413M22 on chromosome 11,
complete sequence.
ACCESSION AL772234
VERSION AL772234
KEYWORDS AL772234.5 GI:23476661
SOURCE HTG.
ORGANISM Mus musculus (house mouse)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
Pearce, A.
Direct Submission
Submitted (01-OCT-2002) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Oct 2, 2002 this sequence version replaced gi:23337380.
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk

During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. The following
abbreviations are used to associate primary accession numbers given
in the feature table with their source databases: Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP
database can be found at
http://www.sanger.ac.uk/Projects/C.elegans/wormpep RP23-413M22 is
from the RPCI-23 Mouse PAC Library
constructed by the group of Pieter de Jong.
For further details see http://www.chori.org/bacpac/home.htm
VECTOR: pBACE3.6.
Location/Qualifiers
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/chromosome="11"
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FEATURES
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ORIGIN
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Best Local Similarity 90.9%; Pred. No. 1.7e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: August 12, 2005, 22:21:57 ; Search time 184.245 Seconds
(without alignments)
738.985 Million cell updates/sec

Title: US-10-673-854-2

Perfect score: 23

Sequence: 1 ccttttcacccactaccactga 23

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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- 1: Geneseqn1980s:*
- 2: Geneseqn1990s:*
- 3: Geneseqn2000s:*
- 4: Geneseqn2001as:*
- 5: Geneseqn2001bs:*
- 6: Geneseqn2002as:*
- 7: Geneseqn2002bs:*
- 8: Geneseqn2003as:*
- 9: Geneseqn2003bs:*
- 10: Geneseqn2003cs:*
- 11: Geneseqn2003ds:*
- 12: Geneseqn2004as:*
- 13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	18.2	79.1	3001	3 AAH51779	Aah51779 Chromosom
C 2	18.2	79.1	53242	11 ACN43926	Acn43926 Human gen
C 3	17.8	77.4	110000	12 AD079173_0	Ado79173 KLF12 gen
C 4	17.4	75.7	50287	11 ACN44848	Acn44848 Mouse gen
C 5	17.4	75.7	81684	13 ABD33502	Abd33502 Murine ca
C 6	17.2	74.8	377	3 AAC28339	Aac28339 Human sec
C 7	17.2	74.8	439	8 ABX38985	Abx38985 Bovine ES
C 8	17.2	74.8	447	13 ADS59130	Ads59130 Bacterial
C 9	17.2	74.8	512	8 ABZ36846	Abz36846 Human GEN
C 10	17.2	74.8	588	12 ACH76108	Ach76108 Human gen
C 11	17.2	74.8	661	6 ABQ99614	Abq99614 Human mem
C 12	17.2	74.8	728	3 AAC93484	Aac93484 Human sec
C 13	17.2	74.8	728	8 ABZ73565	Abz73565 Secretd
C 14	17.2	74.8	728	8 ADA98080	Ada98080 Human sec
C 15	17.2	74.8	728	10 ABZ67162	Abz67162 Human sec
C 16	17.2	74.8	807	12 ACH89809	Ach89809 Human gen
C 17	17.2	74.8	1107	5 AAS70101	Aas70101 DNA encod
C 18	17.2	74.8	1164	9 AAL56877	Aal56877 DNA encod
C 19	17.2	74.8	1194	9 AAL56878	Aal56878 Human ade
C 20	17.2	74.8	1274	8 ABZ74428	Abz74428 Secretd

21	17.2	74.8	1274	8 ADA98866	Ada98866 Human sec
22	17.2	74.8	1374	10 ABZ67985	Abz67985 Human sec
23	17.2	74.8	1402	2 AAQ12228	Aaq12228 Aq41 shor
24	17.2	74.8	1402	2 AAQ24432	Aaq24432 Tak short
25	17.2	74.8	1939	2 AAQ12227	Aaq12227 Aq41 long
26	17.2	74.8	1939	2 AAQ24431	Aaq24431 Tak long
C 27	17.2	74.8	2061	8 ABX71138	Abx71138 Novel hum
C 28	17.2	74.8	3076	2 AAV73003	Aav73003 Human adu
C 29	17.2	74.8	3076	6 ABQ92028	Abq92028 Human pol
C 30	17.2	74.8	3544	6 ABA01937	Abao1937 Human qui
C 31	17.2	74.8	4731	8 ACA45015	ACA45015 Prokaryot
C 32	17.2	74.8	4770	10 ADF00541	Adf00541 Bacterial
C 33	17.2	74.8	11866	1 AAN40141	Aan40141 Sequence
C 34	17.2	74.8	11873	1 AAN40176	Aan40176 Sequence
C 35	17.2	74.8	38736	6 ABQ99652	Abq99652 Human mem
C 36	17.2	74.8	57728	4 AAC87588	Aac87588 Human 9pl
C 37	17.2	74.8	162450	3 AAZ86967	Aaz86967 Retinobla
C 38	16.8	73.0	128	4 AAI27140	Aai27140 Probe #17
C 39	16.8	73.0	128	4 ABA75421	Aba75421 Human foe
C 40	16.8	73.0	128	4 AAI55992	Aai55992 Probe #24
C 41	16.8	73.0	128	4 ABA40051	Abao40051 Probe #18
C 42	16.8	73.0	128	4 AAK50052	Aak50052 Human bon
C 43	16.8	73.0	128	4 AAK23986	Aak23986 Human bra
C 44	16.8	73.0	128	4 ABS49694	Abs49694 Human liv
C 45	16.8	73.0	396	5 AAH65658	Aah65658 C glutami

ALIGNMENTS

RESULT 1

AAH51779/c

ID AAH51779 standard; DNA; 3001 BP.

XX AC AAH51779;

XX DT 29-AUG-2001 (first entry)

XX DE Chromosome 13q31-q33 biallelic marker containing amplicon SEQ ID 191.

XX KW sbg1; g34665; sbg2; g35017; g35018; chromosome 13q31-q33; haplotype;
XX KW biallelic marker; polymorphism; schizophrenia; bipolar disorder; ds.
XX OS Homo sapiens.
XX PN WO200058510-A2.
XX PD 05-OCT-2000.
XX PF 30-MAR-2000; 2000WO-IB000435.
XX PR 30-MAR-1999; 99US-0126903P.
XX PR 30-APR-1999; 99US-0131971P.
XX PR 30-APR-1999; 99US-0132065P.
XX PR 14-JUL-1999; 99US-0143928P.
XX PR 27-JUL-1999; 99US-0145915P.
XX PR 29-JUL-1999; 99US-0146452P.
XX PR 29-JUL-1999; 99US-0146453P.
XX PR 28-OCT-1999; 99US-0162288P.
(GEST) GENSET.

PI Cohen D, Blumenfeld M, Chumakov I, Bougueleret L, Bihain B;
PI Essioux L;
PI WPI; 2000-619082/59.

PT Polynucleotides comprising sequences from sbg1 and g35018 biallelic
PT markers are used for genotyping and detecting schizophrenia or bipolar
PT disorder and predisposition to these disorders.

PS Claim 2; Page 684-685; 737pp; English.

XX

CC AAH51601 represents a human genomic nucleotide sequence comprising sbg1,
 CC g34665, sbg2, g35017 and g35018 nucleic acid sequences located on the
 CC human chromosome 13q31-q33 locus. The nucleotide sequences contain
 CC biallelic markers and polymorphisms. Sequences AAH51602 - AAH51626 and
 CC AAB62907 - AAB62915 represent cDNA human sbg1 cDNA sequences and protein
 CC products. AAH51627 - AAH51631 and AAB62916 - AAB62918 represent g35018
 CC cDNA sequences and protein products. Primers AAH51632 - AAH51699 are used
 CC to isolate sbg1 cDNAs, while sbg1 exons from different primates are
 CC represented by sequences AAH51642 - AAH51699. Nucleotide sequences of
 CC amplicons which comprise biallelic markers located on the chromosome
 CC 13q31-q33 locus are represented in AAH51700 - AAH51817. Biallelic markers
 CC are represented in the sequences by degenerate/undefined base codes. PCR
 CC primers AAH51818 and AAH51819 are used in the isolation of sequences of
 CC the invention. The biallelic marker containing nucleotide sequences are
 CC used to determine the identity of the nucleotide at a biallelic marker in
 CC a sample DNA sequence. The nucleotide sequences may be labelled and used
 CC for genotyping by determining the identity of a nucleotide at a Region D-
 CC related biallelic marker in a biological sample from single or multiple
 CC subjects. By determining the frequency of a biallelic marker in a
 CC population an association between a genotype and a trait, a haplotype and
 CC a trait and a phenotype and a trait can be detected. The sequences can be
 CC used to determine a predisposition to or early onset of schizophrenia or
 CC bipolar disorder or a beneficial response to or side effects related to
 CC treatment against schizophrenia or bipolar disorder

SQ Sequence 3001 BP; 952 A; 600 C; 485 G; 963 T; 0 U; 1 Other;

Query Match 79.1%; Score 18.2; DB 3; Length 3001;
 Best Local Similarity 87.0%; Pred. No. 2e+02;
 Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCCACTACCACTGA 23

Db 1061 CCTGTTCACTCACTACCACTAA 1039

RESULT 2
 ACN43926/c
 ID ACN43926 standard; DNA; 53242 BP.

XX ACN43926;

DT 18-NOV-2004 (first entry)

DE Human genomic sequence HCG1782215.

XX Cytostatic; carcinoma; lymphoma; cancer; human; gene; ss.

XX Homo sapiens.

XX WO2003073826-A2.

PD 12-SEP-2003.

XX 28-FEB-2003; 2003WO-US006235.

XX 01-MAR-2002; 2002US-00087192.

XX (SAGR-) SAGRES DISCOVERY.

XX Morris DW;

XX WPI; 2003-328604/31.

XX Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
 PT comprises a nucleotide sequence.

XX Claim 1; SEQ ID NO 118; Opp; English.

XX The present invention relates to novel DNA and protein sequences which
 CC are associated with carcinomas. The sequences are useful for: (i) for
 CC screening drug candidates; (ii) for screening of bioactive agent capable
 CC of binding to Carcinoma Associated Protein (CAP); (iii) for screening of

CC a bioactive agent capable of modulating the activity of CAP; (iv) for
 CC evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
 CC carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
 CC carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
 CC (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
 CC determining Carcinoma Associated (CA) gene copy number. In addition, the
 CC CA genes are useful as DNA vaccines and the CAP are useful as markers of
 CC carcinoma including lymphoma. The present sequence is one such CA coding
 CC sequence. Note: This patent is an equivalent to basic patent
 CC US2002182586A1, for which no sequence data was published

SQ Sequence 53242 BP; 12636 A; 10850 C; 11247 G; 15524 T; 0 U; 2995 Other;

Query Match 79.1%; Score 18.2; DB 11; Length 53242;

Best Local Similarity 87.0%; Pred. No. 2.7e+02;

Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCCACTACCACTGA 23

Db 10476 CCTTTTCATCCCACTACCACTGA 10454

RESULT 3
 ADO79173_0
 WP Sequence split into 5 fragments LOCUS ADO79173 Accession ADO79173

WP Fragment Name Begin End

WP ADO79173_0 1 110000

WP ADO79173_1 100001 210000

WP ADO79173_2 200001 310000

WP ADO79173_3 300001 410000

WP ADO79173_4 400001 447894

ID ADO79173 standard; DNA; 447894 BP.

XX AC ADO79173;

XX 26-AUG-2004 (first entry)

XX KLF12 gene and surrounding region, SEQ ID 1.

DE Cytostatic; Gene Therapy; human; breast cancer;

XX zinc finger transcriptional repressor; Kruppel-like factor 12; KLF12;

XX transcription factor; AP-2 alpha; AP-2a; chromosome 13; gene; ds;

XX single nucleotide polymorphism; SNP.

XX Homo sapiens.

XX Key Location/Qualifiers

FT variation 1

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FT /standard_name= "Biallelic marker"

FT 1437

FT /tag= b

FT /standard_name= "Biallelic marker"

FT 1604

FT /tag= c

FT /standard_name= "Biallelic marker"

FT 2251

FT /tag= d

FT /standard_name= "Biallelic marker"

FT 2906

FT /tag= e

FT /standard_name= "Biallelic marker"

FT 2953

FT /tag= f

FT /standard_name= "Biallelic marker"

FT 3131

FT /tag= g

FT /standard_name= "Biallelic marker"

FT 3133

FT /tag= h

FT /standard_name= "Biallelic marker"

FT 3238

FT /tag= i

FT /standard_name= "Biallelic marker"

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FT		associated allele is G"	FT		/tag= ak
FT	variation	/tag= w	FT		/standard_name= "Single nucleotide polymorphism"
FT		/standard_name= "Single nucleotide polymorphism"	FT		/note= "This SNP is described as a C/A SNP and the cancer
FT		/note= "This SNP is described as a T/G SNP and the cancer	FT	variation	96429
FT		associated allele is T"	FT		/tag= al
FT	variation	/tag= x	FT		/standard_name= "Single nucleotide polymorphism"
FT		/standard_name= "Single nucleotide polymorphism"	FT		/note= "This SNP is described as a A/T SNP and the cancer
FT		/note= "This SNP is described as a G/T SNP and the cancer	FT	variation	96535
FT		associated allele is T"	FT		/tag= am
FT	variation	/tag= y	FT		/standard_name= "Single nucleotide polymorphism"
FT		/standard_name= "Single nucleotide polymorphism"	FT		/note= "This SNP is described as a C/T SNP and the cancer
FT		/note= "This SNP is described as a G/A SNP and the cancer	FT	variation	110920
FT		associated allele is A"	FT		/tag= an
FT		78067	FT		/standard_name= "Single nucleotide polymorphism"
FT	variation	/tag= z	FT		/note= "This SNP is described as a A/C SNP and the cancer
FT		/standard_name= "Single nucleotide polymorphism"	FT		associated allele is C"
FT		/note= "This SNP is described as a G/A SNP and the cancer	FT	variation	114117

```
FT      /*tag= ao
FT      /standard name= "Single nucleotide polymorphism"
FT      /note= "This SNP is described as a T/C SNP and the cancer
FT      associated allele is T"
FT      117297

Query Match      77.4%; Score 17.8; DB 12; Length 110000;
Best Local Similarity 90.5%; Pred. No. 4.4e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      3 TTTTCATCCAACTACCACTGA 23
      ||||| ||||| |||||
Db      52745 TTTTCATCAACTTCCACTGA 52765

RESULT 4
ACN44848/c
ID ACN44848 standard; DNA; 50287 BP.
XX
AC ACN44848;
XX
DT 18-NOV-2004 (first entry)
XX
DE Mouse genomic sequence mCG4799.
XX
KW Cytostatic; carcinoma; lymphoma; cancer; murine; gene; ss.
XX
OS Mus musculus.
XX
PN WO2003073826-A2.
XX
PD 12-SEP-2003.
XX
PF 28-FEB-2003; 2003WO-US006235.
XX
PR 01-MAR-2002; 2002US-00087192.
XX
PA (SAGR-) SAGRES DISCOVERY.
XX
PI Morris DW;
XX
WPI; 2003-328604/31.
XX
Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
comprises a nucleotide sequence.
XX
Claim 1; SEQ ID NO 1501; Opp; English.
XX
The present invention relates to novel DNA and protein sequences which
are associated with carcinomas. The sequences are useful for: (i) for
screening drug candidates; (ii) for screening of bioactive agent capable
of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
a bioactive agent capable of modulating the activity of CAP; (iv) for
evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
(x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
determining Carcinoma Associated (CA) gene copy number. In addition, the
CA genes are useful as DNA vaccines and the CAP are useful as markers of
carcinoma including lymphoma. The present sequence is one such CA coding
sequence. Note: This patent is an equivalent to basic patent
US2002182586A1, for which no sequence data was published
XX
Sequence 50287 BP; 13307 A; 9608 C; 9890 G; 15956 T; 0 U; 1526 Other;

Query Match      75.7%; Score 17.4; DB 11; Length 50287;
Best Local Similarity 94.7%; Pred. No. 6.2e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 CCTTTTCATCCAACTACCA 19
      ||||| ||||| |||||
Db      48726 CCTTTTCTTCCAACTACCA 48708

/*tag= ao
/standard name= "Single nucleotide polymorphism"
/note= "This SNP is described as a T/C SNP and the cancer
associated allele is T"
117297

Query Match      77.4%; Score 17.8; DB 12; Length 110000;
Best Local Similarity 90.5%; Pred. No. 4.4e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      3 TTTTCATCCAACTACCACTGA 23
      ||||| ||||| |||||
Db      52745 TTTTCATCAACTTCCACTGA 52765

RESULT 4
ACN44848/c
ID ACN44848 standard; DNA; 50287 BP.
XX
AC ACN44848;
XX
DT 18-NOV-2004 (first entry)
XX
DE Mouse genomic sequence mCG4799.
XX
KW Cytostatic; carcinoma; lymphoma; cancer; murine; gene; ss.
XX
OS Mus musculus.
XX
PN WO2003073826-A2.
XX
PD 12-SEP-2003.
XX
PF 28-FEB-2003; 2003WO-US006235.
XX
PR 01-MAR-2002; 2002US-00087192.
XX
PA (SAGR-) SAGRES DISCOVERY.
XX
PI Morris DW;
XX
WPI; 2003-328604/31.
XX
Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
comprises a nucleotide sequence.
XX
Claim 1; SEQ ID NO 1501; Opp; English.
XX
The present invention relates to novel DNA and protein sequences which
are associated with carcinomas. The sequences are useful for: (i) for
screening drug candidates; (ii) for screening of bioactive agent capable
of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
a bioactive agent capable of modulating the activity of CAP; (iv) for
evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
(x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
determining Carcinoma Associated (CA) gene copy number. In addition, the
CA genes are useful as DNA vaccines and the CAP are useful as markers of
carcinoma including lymphoma. The present sequence is one such CA coding
sequence. Note: This patent is an equivalent to basic patent
US2002182586A1, for which no sequence data was published
XX
Sequence 50287 BP; 13307 A; 9608 C; 9890 G; 15956 T; 0 U; 1526 Other;

Query Match      75.7%; Score 17.4; DB 11; Length 50287;
Best Local Similarity 94.7%; Pred. No. 6.2e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 CCTTTTCATCCAACTACCA 19
      ||||| ||||| |||||
Db      48726 CCTTTTCTTCCAACTACCA 48708
```

```
RESULT 5
ABD33502
ID ABD33502 standard; DNA; 81684 BP.
XX
AC ABD33502;
XX
DT 18-NOV-2004 (first entry)
XX
DE Murine cancer-associated (CA) gene MD07-098.
XX
KW Mouse; cancer-associated protein; CAP; cancer-associated gene; CA; gene;
ds; cancer; cytostatic.
XX
OS Mus musculus.
XX
PN WO2004058146-A2.
XX
PD 15-JUL-2004.
XX
PF 15-DEC-2003; 2003WO-US040081.
XX
PR 17-DEC-2002; 2002US-00322281.
XX
PA (SAGR-) SAGRES DISCOVERY INC.
XX
PI Morris DW, Malandro MS;
XX
WPI; 2004-499109/47.
XX
Novel human cancer associated protein encoded within open reading frame
of cancer associated gene, useful as targets for diagnosing cancer.
XX
Disclosure; SEQ ID NO 673; 182pp; English.
XX
The invention relates to cancer-associated proteins (CAP) and the cancer-
associated (CA) nucleic acids encoding them. The invention also relates
to a method for treating cancers involving administering to a patient an
inhibitor of CAP, and a method of screening for anticancer activity in a
potential drug involving providing a cell that expresses a CA gene,
contacting a tissue sample derived from a cancer cell with an anticancer
drug candidate and monitoring the effect of the anticancer drug candidate
on expression of the CA gene. The CAP proteins are useful for detecting
cancer associated with expression of a CAP protein in a test cell sample
and for screening for a bioactive agent capable of modulating the
activity of a CAP protein. The CA nucleic acids are useful for diagnosing
cancer, involving determining the expression of a CA nucleic acid in a
tissue. This sequence represents a murine CA gene of the invention. Note:
The sequence data for this patent did not form part of the printed
specification, but was obtained in electronic format directly from WIPO
at ftp.wipo.int/pub/published_pct_sequences
XX
Sequence 81684 BP; 21861 A; 18581 C; 19236 G; 21459 T; 0 U; 547 Other;

Query Match      75.7%; Score 17.4; DB 13; Length 81684;
Best Local Similarity 94.7%; Pred. No. 6.5e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 CCTTTTCATCCAACTACCA 19
      ||||| ||||| |||||
Db      72798 CCTTTTCTTCCAACTACCA 72816

RESULT 6
AAC28339/c
ID AAC28339 standard; cDNA; 377 BP.
XX
AC AAC28339;
XX
DT 06-OCT-2000 (first entry)
XX
DE Human secreted protein 5' EST, SEQ ID NO: 32414.
XX
```

KW Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;
 XX gene therapy; chromosome mapping; ss.

XX Homo sapiens.

XX EP1033401-A2.

XX 06-SEP-2000.

XX 21-FEB-2000; 2000EP-00200610.

XX 26-FEB-1999; 99US-0122487P.

XX (GEST) GENSET.

XX Dumas Milne Edwards J, Duclert A, Giordano J;

XX WPI; 2000-500381/45.

XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for
 PT obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for
 PT diagnostic, forensic, gene therapy and chromosome mapping procedures.

XX Claim 1; SEQ ID NO 32414; 71pp + Sequence Listing; English.

XX The present sequence is one of a large number of 5' ESTs derived from
 CC mRNAs encoding secreted proteins. No ORF has yet been conclusively
 CC identified within the present sequence. The 5' ESTs were prepared from
 CC total human RNAs or polyA+ RNAs derived from 30 different tissues. EST
 CC sequences usually correspond mainly to the 3' untranslated region (UTR)
 CC of the mRNA because they are often obtained from oligo-dT primed cDNA
 CC libraries. Such ESTs are not well suited for isolating cDNA sequences
 CC derived from the 5' ends of mRNAs and even in those cases where longer
 CC cDNA sequences have been obtained, the full 5' UTR is rarely included. 5'
 CC ESTs are derived from mRNAs with intact 5' ends and can therefore be used
 CC to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used in
 CC diagnostic, forensic, gene therapy and chromosome mapping procedures.
 CC They are used to obtain upstream regulatory sequences and to design
 CC expression and secretion vectors

XX SQ Sequence 377 BP; 83 A; 57 C; 73 G; 159 T; 0 U; 5 Other;

Query Match 74.8%; Score 17.2; DB 3; Length 377;

Best Local Similarity 86.4%; Pred. No. 4.8e+02;

Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 CTTTTCATCCAACTACCACTGA 23

Db 288 CATTTCATCCAACTACCACTGA 267

RESULT 7

ABX38985/c
 ID ABX38985 standard; cDNA; 439 BP.

XX AC ABX38985;

XX 20-FEB-2003 (first entry)

XX Bovine EST associated with lactation/muscle/fat deposition #4150.

XX Bovine; ss; EST; expressed sequence tag; lactation; LMPD;

XX muscle deposition; fat deposition; genome mapping; gene identification;

XX gene analysis; cattle breeding.

XX Bos Taurus.

XX US2002137139-A1.

XX 26-SEP-2002.

XX 24-SEP-2001; 2001US-00960352.

PR 12-JAN-1999; 99US-0115707P.
 XX 11-JAN-2000; 2000US-00480902.

XX (BYAT/) BYATT J C.

XX (MATH/) MATHIALAGAN N.

XX (TAON/) TAO N.

XX (WARR/) WARREN W C.

XX Byatt JC, Mathialagan N, Tao N, Warren WC;

XX WPI; 2003-110599/10.

XX New nucleic acid associated with lactation, and muscle and fat
 PT deposition, useful for genome mapping, gene identification and analysis,
 PT cattle breeding, or for genetically improving cattle.

XX Claim 2; SEQ ID NO 4150; 245pp; English.

XX The invention relates to a purified nucleic acid molecule associated with
 CC lactation or muscle and fat deposition (designated LMPD), derived from
 CC cattle, and the LMPD nucleic acid can specifically hybridise to a second
 CC nucleic acid molecule comprising any of 15112 nucleotide sequences,
 CC appearing as ABX34836-ABX49947, or complements of them. Also included are
 CC ; (1) a transformed cell having a nucleic acid comprising an LMPD nucleic
 CC acid linked to a promoter and a 3' non-translated sequence that
 CC functions in the cell to cause termination of transcription and addition
 CC of polyadenylated ribonucleotides to a 3' end of the mRNA molecule; and
 CC (2) determining a level or pattern of a molecule in a bovine cell or
 CC tissue comprising: (a) incubating a marker nucleic acid (comprising any
 CC of the 15112 nucleic acid sequences or its complement or fragment) with a
 CC complementary nucleic acid molecule obtained from the bovine cell or
 CC tissue, where hybridisation between the marker nucleic acid and the
 CC complementary nucleic acid permits the detection of the molecule; and (b)
 CC detecting the level or pattern of the complementary nucleic acid, where
 CC the detection of the complementary nucleic acid is predictive of the
 CC level or pattern of the molecule. The LMPD nucleic acid is used for
 CC determining a level or pattern of a molecule in a bovine cell or tissue.
 CC It is useful for genome mapping, gene identification and analysis, cattle
 CC breeding, preparation of constructs for use in cattle gene expression, or
 CC for genetically improving cattle. The present sequence is one of the
 CC 15112 bovine LMPD EST (expressed sequence tag) nucleic acids. Note: The
 CC present sequence was not shown in the specification but was obtained in
 CC electronic format from the USPTO web site:
 CC seqdata.uspto.gov/sequence.html?DocID=20020137139

XX SQ Sequence 439 BP; 152 A; 58 C; 80 G; 149 T; 0 U; 0 Other;

Query Match 74.8%; Score 17.2; DB 8; Length 439;

Best Local Similarity 86.4%; Pred. No. 4.9e+02;

Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 CTTTTCATCCAACTACCACTGA 23

Db 251 CTTTTCATGAACTACCACTGA 230

RESULT 8

AD559130/c

ID AD559130 standard; cDNA; 447 BP.

XX AC AD559130;

XX 02-DEC-2004 (first entry)

XX Bacterial polynucleotide #11117.

XX Recombinant DNA construct; transformed plant; improved plant property;
 KW cold tolerance; heat tolerance; drought tolerance; herbicide; osmosis;
 KW pathogen tolerance; pest tolerance; plant disease resistance;
 KW cell cycle pathway modification; plant growth regulator;
 KW homologous recombination; seed oil yield; protein yield; carbohydrate;
 KW nitrogen; phosphorus; photosynthesis; lignin; galactomannan;
 KW bacterial polynucleotide; gene; ss.

PA (RANK/) RANK D R.
XX (HANKZ/) HANKZEL D K.
PI Penn SG, Rank DR, Hanzel DR;
XX WPI; 2004-119264/12.
XX
XX New human genome-derived single exon nucleic acid probes useful for human
PT gene expression analysis, for identifying or characterizing alternative
PT splicing events, for assessing genomic alterations or as tools for
PT surveying tissues.
XX
XX Claim 15; SEQ ID NO 9303; 80pp; English.
XX
XX The invention relates to a nucleic acid probe for measuring human gene
CC expression, comprising any of the 27,400 fully defined nucleotide
CC sequences in the specification, or their complements or fragments, and
CC encoding at least 8 amino acids of any of the 688 amino acid sequences
CC fully defined in the specification. The probe is a single exon probe that
CC hybridises under high stringency conditions to a nucleic acid molecule
CC expressed in human cells or tissues. Also included are a spatially-
CC addressable set of single exon nucleic acid probes for measuring human
CC gene expression (comprising a plurality of single exon nucleic acid
CC probes cited above, where each of the plurality of probes is separately
CC and addressably isolatable or amplifiable from the plurality), a single
CC exon microarray for measuring human gene expression, a method of
CC measuring human gene expression, a vector comprising the single exon
CC probe cited above, an ORF-encoded peptide comprising at least 8
CC contiguous amino acids of any of the above-mentioned amino acid
CC sequences (optionally with conservative amino acid substitutions), an
CC isolated antibody that binds specifically to a peptide cited above,
CC methods of selling and/or licensing single exon probes or microarrays to
CC a customer desiring to measure gene expression, a method of providing
CC human gene expression data by subscription, and a computer-readable
CC storage medium which contains a database having a plurality of records
CC (each record including data on the expression of a single exon probe
CC cited above. The probe, methods and apparatus are useful in gene
CC expression analysis. The probes may be used as tools for surveying
CC tissues to detect the presence of expressed messages that contain their
CC specific exon, or in constructing genome-derived single exon microarrays.
CC In addition, the probes are used in identifying and characterising
CC alternative splicing events, in detecting and characterising gross
CC alterations in the genomic locus that includes their exon, in assessing
CC smaller genomic alterations, in priming the synthesis of nucleic acids,
CC or in expressing the ORF-encoded peptide. The present sequence is a human
CC single exon probe of the invention. Note: The sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from USPTO at
CC seqdata.uspto.gov/sequence.html?DocID=20030194704
XX
XX Sequence 588 BP; 253 A; 120 C; 91 G; 124 T; 0 U; 0 Other;
Query Match 74.8%; Score 17.2; DB 12; Length 588;
Best Local Similarity 86.4%; Pred. No. 5e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 2 CTTTTCATCCCACTACCACTGA 23
DB 245 CATTTTCATCCCACTACCACTG 266
RESULT 11
ABQ99614/c
ID ABQ99614 standard; cDNA; 661 BP.
XX
XX ABQ99614;
XX
XX 12-NOV-2002 (first entry)
XX Human membrane spanning 4-domain family, subfamily A 6E cDNA.
XX Human; membrane spanning 4-domain A; cytostatic; antiallergic; MS4A;
KW gene therapy; atopic disorder; non-Hodgkin's lymphoma;
XX

KW Hodgkin's lymphoma; allergenic disease; MS4A6E; gene; ss;
KW chromosome 11q12-13.1.
XX
XX Homo sapiens.
OS
XX
XX Key Location/Qualifiers
FH CDS 15..4586
FT /*tag= a
FT /product= "MS4A6E"
XX
XX WO200262946-A2.
XX
XX 15-AUG-2002.
XX
XX 10-DEC-2001; 2001WO-US048437.
XX
XX 08-DEC-2000; 2000US-0254362P.
XX 20-FEB-2001; 2001US-0270057P.
XX (UYDU-) UNIV DUKE.
XX
XX Tedder TF, Liang YH;
XX
XX WPI; 2002-657530/70.
DR P-PSDB; ABP65028.
XX
XX New membrane spanning 4-domain A (MS4A) genes and polypeptides, useful
PT for generating animal models of atopic disorders, for drug screening, or
PT for treating (non-) Hodgkin's lymphoma, or allergenic or atopic disorders
PT in e.g. humans.
XX
XX Claim 2; Page 125-126; 450pp; English.
XX
XX The invention relates to novel membrane spanning 4-domain A (MS4A)
CC nucleic acid and polypeptide molecules, comprising human and mouse MS4A.
CC The polypeptides of the invention have cytostatic and antiallergic
CC activity. The polynucleotides may have a use in gene therapy. The MS4A
CC nucleic acids and polypeptides are useful for generating animal (e.g.
CC mouse) models of atopic disorders, or for drug discovery screens. These
CC are also useful for treating (non-)Hodgkin's lymphoma, allergenic
CC diseases, atopic disorders or other MS4A-related conditions. The present
CC sequence encodes human membrane spanning 4-domain A 6E (MS4A6E)
XX
XX Sequence 661 BP; 195 A; 147 C; 132 G; 187 T; 0 U; 0 Other;
Query Match 74.8%; Score 17.2; DB 6; Length 661;
Best Local Similarity 86.4%; Pred. No. 5e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 CCTTTTCATCCCACTACCACTG 22
DB 285 CCTTTTCATCCCACTACCTG 264
RESULT 12
AAC93484
ID AAC93484 standard; cDNA; 728 BP.
XX
XX AAC93484;
XX
XX 16-FEB-2001 (first entry)
XX
XX Human secreted protein gene 6 SEQ ID NO:16.
XX
XX Human; secreted protein; immunosuppressive; antiarthritic; antirheumatic;
KW antiproliferative; cytostatic; cardiac; vasotropic; cerebroprotective;
KW notropic; neuroprotective; antibacterial; virucide; fungicide;
KW ophthalmological; vulnery; autoimmune disease; rheumatoid arthritis;
KW hyperproliferative disorders; cancer; cardiovascular disorder;
KW cardiac arrest; cerebrovascular disorder; nervous system disorder;
KW Alzheimer's disease; ocular disorder; wound healing; skin aging; ss.
XX
XX Homo sapiens.
OS

XX WO200061626-A1.
 PN 19-OCT-2000.
 XX 06-APR-2000; 2000WO-US009066.
 XX 09-APR-1999; 99US-0128698P.
 PR 20-JAN-2000; 2000US-0176926P.
 XX (HUMA-) HUMAN GENOME SCI INC.
 PA (ROSE/) ROSEN C A.
 XX Rosen CA, Ruben SM, Komatsoulis G;
 PI WPI; 2000-619227/59.
 XX P-PSDB; AAB51832.
 DR New nucleic acid molecules encoding 49 human secreted proteins for
 PT diagnosing, preventing or ameliorating medical conditions and used for
 PT food additives or preservatives.
 XX Claim 1; Page 435-436; 516pp; English.
 PS Polynucleotide sequences AAC93479 - AAC93527 represent cDNA encoding
 CC human secreted proteins AAB51827 - AAB51875. Sequences AAB51876 -
 CC AAB51927 represent alternative polypeptides encoded by the genes, and
 CC amino acid sequences with which they share homology. The genes and
 CC proteins have activities dependent on the tissues and cells in which they
 CC are expressed. Examples of their activities include immunosuppressive;
 CC antiarthritic; antirheumatic; antiproliferative; cytostatic; cardiac;
 CC vasotropic; cerebroprotective; neurotropic; neuroprotective; antibacterial;
 CC virucide; fungicide; opthalmological; and vulnerary. The secreted
 CC proteins, polynucleotides, antagonists and agonists may be useful in
 CC treating, preventing and/or diagnosing diseases and disorders such as
 CC autoimmune diseases e.g. rheumatoid arthritis, hyperproliferative
 CC disorders e.g. neoplasms of the breast or liver, cardiovascular disorders
 CC e.g. cardiac arrest, cerebrovascular disorders e.g. cerebral ischaemia,
 CC angioneogenesis, nervous system disorders e.g. Alzheimer's disease,
 CC infections caused by bacteria, viruses and fungi and ocular disorders
 CC e.g. corneal infection. The polypeptides can also be used to aid wound
 CC healing and epithelial cell proliferation, to prevent skin aging due to
 CC sunburn, to maintain organs before transplantation, for supporting cell
 CC culture of primary tissues, to regenerate tissues and in chemotaxis. The
 CC polypeptides can also be used as a food additive or preservative to
 CC increase or decrease storage capabilities, fat content, lipid, protein,
 CC carbohydrate, vitamins, minerals, cofactors and other nutritional
 CC components. Oligonucleotides AAC93470 - AAC93478 and peptide AAB51826 are
 CC used in the isolation and characterisation of the proteins and
 CC polynucleotides of the invention
 XX SQ Sequence 728 BP; 178 A; 169 C; 183 G; 198 T; 0 U; 0 Other;
 Query Match 74.8%; Score 17.2; DB 3; Length 728;
 Best Local Similarity 86.4%; Pred. No. 5.1e+02;
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 QY 1 CCTTTTCATCCCACTTACCCTG 22
 ||||| ||||| ||||| ||||| |||||
 Db 178 CCTGTGTCAGCCAACTTCCACTG 199
 RESULT 13
 ABZ73565
 ID ABZ73565 standard; cDNA; 728 BP.
 XX AC ABZ73565;
 XX DT 12-MAY-2003 (first entry)
 XX DE Secreted protein-encoding gene 285 cDNA clone HRDFK37, SEQ ID NO:295.
 XX Human; secreted protein; cancer; tumour; hyperproliferative disorder;
 KW

KW autoimmune disorder; inflammation; angiogenic diseases; AIDS;
 KW acquired immunodeficiency syndrome; hepatitis; anaemia; wound healing;
 KW drug screening; chromosome identification; chromosome mapping;
 KW cytostatic; gene therapy; antiinflammatory; immunomodulator; anti-HIV;
 KW antianaemic; vulnerary; gene; ss.
 XX Homo sapiens.
 XX WO200277013-A2.
 PN 03-OCT-2002.
 PD 26-MAR-2002; 2002WO-US009370.
 PF 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX (HUMA-) HUMAN GENOME SCI INC.
 PA Rosen CA, Ruben SM;
 PI WPI; 2003-040578/03.
 XX P-PSDB; ABR01231.
 DR New human secreted proteins and nucleic acids, useful for detecting or
 PT treating cancer or other hyperproliferative disorders, autoimmune
 PT disorders, inflammatory disorders, HIV disease, hepatitis or anemia.
 XX Claim 21; Page 1307; 2474pp; English.
 PS ABZ73281-ABZ73697 represent cDNAs corresponding to 391 human secreted
 CC protein genes, and ABP00947-ABP01363 represent the proteins they encode.
 CC ABZ73698-ABZ74687 represent human secreted protein genomic fragments. The
 CC invention also encompasses antibodies specific for the secreted proteins,
 CC the use of the secreted proteins in drug screening and recombinant
 CC vectors and host cells comprising a nucleic acid of the invention. The
 CC secreted proteins are thought to be involved in biological activities
 CC associated with cellular signalling, cellular differentiation, cell
 CC migration, prohormone activation and neurotransmitter activity. The
 CC secreted proteins, nucleic acids encoding them, antibodies or antibody
 CC fragments specific for the secreted proteins, and modulators of protein
 CC activity are useful for diagnosing or treating cancers or other
 CC hyperproliferative disorders. Additionally, the secreted proteins and
 CC their nucleic acids may also be used in the treatment of autoimmune
 CC disorders, inflammatory disorders, diseases involving angiogenesis, AIDS
 CC (acquired immunodeficiency syndrome), hepatitis, anaemia, and to promote
 CC wound healing. Nucleic acids of the invention may be used for chromosome
 CC identification, chromosome mapping, in gene therapy, for identifying
 CC individuals from minute biological samples, as hybridisation probes, and
 CC as molecular weight markers. The present sequence represents a human
 CC secreted protein-encoding cDNA clone of the invention
 XX SQ Sequence 728 BP; 178 A; 169 C; 183 G; 198 T; 0 U; 0 Other;
 Query Match 74.8%; Score 17.2; DB 8; Length 728;
 Best Local Similarity 86.4%; Pred. No. 5.1e+02;
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 QY 1 CCTTTTCATCCCACTTACCCTG 22
 ||||| ||||| ||||| ||||| |||||
 Db 178 CCTGTGTCAGCCAACTTCCACTG 199
 RESULT 14
 ADA98080
 ID ADA98080 standard; cDNA; 728 BP.
 XX AC ADA98080;
 XX DT 20-NOV-2003 (first entry)
 XX DE Human secreted protein cDNA sequence #174.

XX human; secreted protein; cardiovascular disorder; arrhythmia;
 KW atherosclerosis; stroke; endocarditis; congestive heart failure;
 KW rheumatic heart disease; cardiomyopathy; haemorrhoids; varicose veins;
 KW migraine; thrombosis; neural disorder; immune system disorder;
 KW muscular disorder; reproductive disorder; gastrointestinal disorder;
 KW pulmonary disorder; renal disorder; proliferative disorder; cancer; gene;
 KW ss.

XX Homo sapiens.
 OS
 XX WO2003004623-A2.
 PN
 XX
 XX 16-JAN-2003.
 PD
 XX
 XX 26-MAR-2002; 2002WO-US009922.
 PF
 XX
 XX 27-MAR-2001; 2001US-0278650P.
 PR
 XX 12-SEP-2001; 2001US-00950082.
 PR
 XX 12-SEP-2001; 2001US-00950083.
 PR
 XX
 XX (HUMA-) HUMAN GENOME SCI INC.
 PA
 XX
 XX Rosen CA, Ruben SM;
 PI
 XX WPI; 2003-247946/24.
 DR
 XX

XX New human secreted polypeptide and nucleic acid molecules, useful for
 PT diagnosing, preventing, prognosticating or treating cardiovascular
 PT disorders (e.g. arrhythmia, atherosclerosis, cardiomyopathy, or
 PT thrombosis).
 XX
 XX Claim 1; SEQ ID NO 184; 1572pp; English.

XX The invention comprises the amino acid and coding sequence of human
 CC secreted proteins. The DNA and protein sequences of the invention are
 CC useful in the treatment of cardiovascular disorders, such as: arrhythmia,
 CC atherosclerosis, stroke, endocarditis, congestive heart failure,
 CC rheumatic heart disease, cardiomyopathy, haemorrhoids, varicose veins,
 CC migraine, or thrombosis. The DNA and protein sequences may also be used
 CC for treating or preventing: neural disorders, immune system disorders,
 CC muscular disorders, reproductive disorders, gastrointestinal disorders,
 CC pulmonary disorders, renal disorders, proliferative disorders and/or
 CC cancerous diseases. The present cDNA sequence encodes a human secreted
 CC protein of the invention. NOTE: The present sequence is shown on the WIPO
 CC website.

XX Sequence 728 BP; 178 A; 169 C; 183 G; 198 T; 0 U; 0 Other;
 SQ
 Query Match 74.8%; Score 17.2; DB 8; Length 728;
 Best Local Similarity 86.4%; Pred. No. 5.1e+02;
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCAACTACCACTG 22
 |||||
 Db 178 CCTTGTGAGCCAACTTCCACTG 199

RESULT 15
 ABZ67162
 ID ABZ67162 standard; cDNA; 728 BP.

XX ABZ67162;

XX 26-MAR-2003 (first entry)

XX Human secreted protein encoding cDNA SEQ ID NO 282.

XX Human; secreted protein; neurotropic; neuroprotective; cytostatic;
 KW virucide; dermatological; immunosuppressive; antiinflammatory; anti-HIV;
 KW vulnery; antibacterial; antiparkinsonian; antisickling; antianaemic;
 KW antiarthritic; cancer; antirheumatic; hepatotropic; cerebroprotective;
 KW antiinflammatory; antiallergic; antidiabetic; antiulcer; anticonvulsant;

KW antifungal; antiparasitic; cardiant; immune disorder; infection; vaccine;
 KW cardiovascular disorder; neurological disease; nephrotropic;
 KW gene therapy; gene; ds.

XX Homo sapiens.
 OS
 XX WO200277186-A2.
 PN
 XX
 XX 03-OCT-2002.
 PD
 XX
 XX 26-MAR-2002; 2002WO-US009188.
 PF
 XX
 XX 27-MAR-2001; 2001US-0278650P.
 PR
 XX 12-SEP-2001; 2001US-00950082.
 PR
 XX 12-SEP-2001; 2001US-00950083.
 PR
 XX
 XX (HUMA-) HUMAN GENOME SCI INC.

XX Rosen CA, Ruben SM;
 PI
 XX WPI; 2003-040583/03.
 DR
 XX P-PSDB; ABP99741.
 DR
 XX

XX New human secreted proteins encoded by genes contained in cDNA clones
 PT (e.g. HGCAC19), useful for preventing, treating or diagnosing e.g. AIDS,
 PT multiple sclerosis, herpes virus, leukemia, tick-borne encephalitis or
 PT West Nile fever.

XX Claim 7; Page 1316; 2423pp; English.

XX The invention relates to novel human genes (ABZ66891-ABZ68209) and the
 CC encoded secreted proteins (ABP99470-ABP99872) useful for preventing,
 CC treating or ameliorating medical conditions e.g. by protein or gene
 CC therapy. The genes are isolated from a range of human tissues disclosed
 CC in the specification. The nucleic acids, proteins, antibodies and
 CC (ant)agonists are useful in the diagnosis, treatment and prevention of:
 CC (a) cancer, e.g. breast and ovarian cancer and other cancers of the
 CC adrenal gland, bone, bone marrow, breast, gastrointestinal tract, liver,
 CC lung or urogenital; (b) immune disorders e.g. Addison's disease,
 CC allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis,
 CC diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid
 CC arthritis and ulcerative colitis; (c) cardiovascular disorders such as
 CC myocardial ischaemia; (d) wound healing; (e) neurological diseases e.g.
 CC cerebral anoxia and epilepsy; and (f) infectious diseases such as viral,
 CC bacterial, fungal and parasitic infections

XX Sequence 728 BP; 178 A; 169 C; 183 G; 198 T; 0 U; 0 Other;

Query Match 74.8%; Score 17.2; DB 10; Length 728;
 Best Local Similarity 86.4%; Pred. No. 5.1e+02;
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCAACTACCACTG 22
 |||||
 Db 178 CCTTGTGAGCCAACTTCCACTG 199

Search completed: August 13, 2005, 04:14:35
 Job time : 191.245 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: August 13, 2005, 03:33:32 ; Search time 56.5213 Seconds
(without alignments)
665.844 Million cell updates/sec

Title: US-10-673-854-2
Perfect score: 23
Sequence: 1 cctttcatccaactaccactga 23

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents NA.*
1: /cgn2_6/ptodata/1/ina/5A COMB.seq.*
2: /cgn2_6/ptodata/1/ina/5B COMB.seq.*
3: /cgn2_6/ptodata/1/ina/6A COMB.seq.*
4: /cgn2_6/ptodata/1/ina/6B COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PCTUS COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	18.2	79.1	3001	4	US-09-539-333D-191
C 2	17.8	77.4	103447	4	US-09-949-016-16320
C 3	17.2	74.8	377	4	US-09-513-999C-32414
C 4	17.2	74.8	513	4	US-09-621-976-1795
C 5	17.2	74.8	562	4	US-09-621-976-17125
C 6	17.2	74.8	601	4	US-09-949-016-96155
C 7	17.2	74.8	601	4	US-09-949-016-96156
C 8	17.2	74.8	2061	4	US-09-774-528-366
C 9	17.2	74.8	4770	4	US-09-543-681A-826
C 10	17.2	74.8	36755	4	US-09-949-016-16994
C 11	17.2	74.8	103750	4	US-09-949-016-13319
C 12	17.2	74.8	162450	3	US-09-345-882-1
C 13	17.2	74.8	387902	4	US-09-949-016-14543
C 14	17.2	74.8	421883	4	US-09-949-016-12557
C 15	16.8	73.0	942	4	US-09-328-352-3203
C 16	16.8	73.0	15478	4	US-09-949-016-15288
C 17	16.8	73.0	74644	4	US-09-949-016-17556
C 18	16.8	73.0	784019	4	US-09-949-016-14033
C 19	16.8	73.0	828152	4	US-09-949-016-12777
C 20	16.6	72.2	601	4	US-09-949-016-199769
C 21	16.6	72.2	601	4	US-09-949-016-199770
C 22	16.6	72.2	601	4	US-09-949-016-199771
C 23	16.6	72.2	325791	4	US-09-768-185A-1
C 24	16.6	72.2	373182	4	US-09-949-016-17371
C 25	16.6	72.2	373694	4	US-09-949-016-12062
C 26	16.4	71.3	601	4	US-09-949-016-169822
C 27	16.4	71.3	47115	4	US-09-949-016-12278

C 28	16.4	71.3	47122	4	US-09-949-016-16520	Sequence 16520, A
C 29	16.2	70.4	282	4	US-09-107-433-487	Sequence 487, App
C 30	16.2	70.4	298	4	US-09-513-999C-36033	Sequence 36033, A
C 31	16.2	70.4	482	4	US-09-495-050A-114	Sequence 114, App
C 32	16.2	70.4	528	4	US-09-107-433-1306	Sequence 1306, App
C 33	16.2	70.4	601	4	US-09-949-016-42842	Sequence 42842, A
C 34	16.2	70.4	601	4	US-09-949-016-119957	Sequence 119957, A
C 35	16.2	70.4	601	4	US-09-949-016-133993	Sequence 133993, A
C 36	16.2	70.4	633	4	US-09-248-796A-906	Sequence 906, App
C 37	16.2	70.4	869	3	US-08-998-416-491	Sequence 491, App
C 38	16.2	70.4	921	4	US-09-583-110-47	Sequence 47, Appli
C 39	16.2	70.4	1395	3	US-08-834-776A-1	Sequence 1, Appli
C 40	16.2	70.4	1887	4	US-09-107-532A-3469	Sequence 3469, Ap
C 41	16.2	70.4	1941	4	US-09-248-796A-2598	Sequence 2598, Ap
C 42	16.2	70.4	2092	4	US-09-545-773-3	Sequence 3, Appli
C 43	16.2	70.4	2148	4	US-09-949-016-2145	Sequence 2145, Ap
C 44	16.2	70.4	7101	1	US-08-480-604A-9	Sequence 9, Appli
C 45	16.2	70.4	7101	2	US-08-405-496A-9	Sequence 9, Appli

RESULT 1
US-09-539-333D-191/c
; Sequence 191, Application US/09539333D
; Patent No. 6476208
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; APPLICANT: Bougueleret, Lydie
; APPLICANT: Bihain, Bernard
; APPLICANT: Essioux, Laurent
; TITLE OF INVENTION: SCHIZOPHRENIA ASSOCIATED GENES, PROTEINS AND BIALLELIC MARKERS
; FILE REFERENCE: GENSET.047AUS
; CURRENT APPLICATION NUMBER: US/09/539,333D
; CURRENT FILING DATE: 2000-03-30
; PRIOR APPLICATION NUMBER: US 60/126,903
; PRIOR FILING DATE: 1999-03-30
; PRIOR APPLICATION NUMBER: US 60/131,971
; PRIOR FILING DATE: 1999-04-30
; PRIOR APPLICATION NUMBER: US 60/132,065
; PRIOR FILING DATE: 1999-04-30
; PRIOR APPLICATION NUMBER: US 60/143,928
; PRIOR FILING DATE: 1999-07-14
; PRIOR APPLICATION NUMBER: US 60/145,915
; PRIOR FILING DATE: 1999-07-27
; PRIOR APPLICATION NUMBER: US 60/146,453
; PRIOR FILING DATE: 1999-07-29
; PRIOR APPLICATION NUMBER: US 60/146,452
; PRIOR FILING DATE: 1999-07-29
; PRIOR APPLICATION NUMBER: US 60/162,288
; PRIOR FILING DATE: 1999-10-28
; PRIOR APPLICATION NUMBER: US 09/416,384
; PRIOR FILING DATE: 1999-10-12
; NUMBER OF SEQ ID NOS: 231
; SOFTWARE: Patent.pm
; SEQ ID NO 191
; LENGTH: 3001
; TYPE: DNA
; ORGANISM: Homo Sapiens
; FEATURE:
; NAME/KEY: allele
; LOCATION: 1501
; OTHER INFORMATION: 99-26191-58 : polymorphic base G or A
; FEATURE:
; NAME/KEY: misc binding
; LOCATION: 1502..1520
; OTHER INFORMATION: 99-26191-58.misl, complement
; FEATURE:
; NAME/KEY: misc binding
; LOCATION: 1481..1500

ALIGNMENTS

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; OTHER INFORMATION: 99-26191-58.mis2,
; FEATURE:
; NAME/KEY: primer_bind
; LOCATION: 1539..1558
; OTHER INFORMATION: upstream amplification primer, complement
; FEATURE:
; NAME/KEY: primer_bind
; LOCATION: 1095..1115
; OTHER INFORMATION: downstream amplification primer
; FEATURE:
; NAME/KEY: misc_binding
; LOCATION: 1489..1513
; OTHER INFORMATION: 99-26191-58 probe
US-09-539-333D-191

Query Match          79.1%; Score 18.2; DB 4; Length 3001;
Best Local Similarity 87.0%; Pred. No. 43;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCCACTACCACTGA 23
Db 1061 CCGTTCATTCACCTACCACTAA 1039

RESULT 2
US-09-949-016-16320/c
; Sequence 16320, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16320
; LENGTH: 103447
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(103447)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-16320

Query Match          77.4%; Score 17.8; DB 4; Length 103447;
Best Local Similarity 90.5%; Pred. No. 11e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TTTTCATCCCACTACCACTGA 23
Db 7931 TTTCCATCAACTACCACTGA 7911

RESULT 3
US-09-513-999C-32414/c
; Sequence 32414, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
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; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 32414
; LENGTH: 377
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 144_feature
; OTHER INFORMATION: r-a or g
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 236
; OTHER INFORMATION: n=a, g, c or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 261
; OTHER INFORMATION: n=a, g, c or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 335
; OTHER INFORMATION: d=a or g or t
US-09-513-999C-32414

Query Match          74.8%; Score 17.2; DB 4; Length 377;
Best Local Similarity 86.4%; Pred. No. 95;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 CTTTTCATCCCACTACCACTGA 23
Db 288 CATTTCATCCCACTACCACTGA 267

RESULT 4
US-09-621-976-1795/c
; Sequence 1795, Application US/09621976
; Patent No. 6639063
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Jobert, S.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: ESTs and Encoded Human Proteins.
; FILE REFERENCE: GENSET.054PR2
; CURRENT APPLICATION NUMBER: US/09/621,976
; CURRENT FILING DATE: 2000-07-21
; NUMBER OF SEQ ID NOS: 19335
; SOFTWARE: Patent.pm
; SEQ ID NO 1795
; LENGTH: 513
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 5..409
; NAME/KEY: sig_peptide
; LOCATION: 5..181
; OTHER INFORMATION: Von Heijne matrix
; OTHER INFORMATION: score 4.5
; OTHER INFORMATION: seq LSALSALVGFILL/SV
US-09-621-976-1795

Query Match          74.8%; Score 17.2; DB 4; Length 513;
Best Local Similarity 86.4%; Pred. No. 99;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
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QY 1 CCTTTTCATCCAACTACCACTG 22
|||||
Db 239 CCTTTTCGTCCTCACTTACACTG 218

RESULT 5

US-09-621-976-17125/c
; Sequence 17125, Application US/09621976
; Patent No. 6639063
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Jobert, S. J.Y.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: ESTs and Encoded Human Proteins.
; FILE REFERENCE: GENSET.054PR2
; CURRENT APPLICATION NUMBER: US/09/621,976
; CURRENT FILING DATE: 2000-07-21
; NUMBER OF SEQ ID NOS: 19335
; SOFTWARE: Patent.pm
; SEQ ID NO 17125
; LENGTH: 562
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-621-976-17125

Query Match 74.8%; Score 17.2; DB 4; Length 562;
Best Local Similarity 86.4%; Pred. No. 1e+02; 3; Indels 0; Gaps 0;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCAACTACCACTG 22
|||||
Db 331 CCTTTTCGTCCTCACTTACACTG 310

RESULT 6

US-09-949-016-96155
; Sequence 96155, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 96155
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-96155

Query Match 74.8%; Score 17.2; DB 4; Length 601;
Best Local Similarity 86.4%; Pred. No. 1e+02; 3; Indels 0; Gaps 0;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CCTTTTCATCCAACTACCACTG 23
|||||
Db 368 CATTTTCACCAACACACACAGA 389

RESULT 7

US-09-949-016-96156
; Sequence 96156, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:

; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 96156
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-96156

Query Match 74.8%; Score 17.2; DB 4; Length 601;
Best Local Similarity 86.4%; Pred. No. 1e+02; 3; Indels 0; Gaps 0;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CCTTTTCATCCAACTACCACTG 23
|||||
Db 438 CATTTTCACCAACACACACAGA 459

RESULT 8

US-09-774-528-366/c
; Sequence 366, Application US/09774528
; Patent No. 6743619
; GENERAL INFORMATION:
; APPLICANT: Tang, Y. Tom
; APPLICANT: Zhou, Ping
; APPLICANT: Goodrich, Ryle
; APPLICANT: Liu, Chenghua
; APPLICANT: Asundi, Vinod
; APPLICANT: Ren, Feiyan
; APPLICANT: Zhang, Jie
; APPLICANT: Zhao, Qing A.
; APPLICANT: Yang, Yonghong
; APPLICANT: Xue, Aidong J.
; APPLICANT: Wehrman, Tom
; APPLICANT: Wang, Jian-Rui
; APPLICANT: Wang, Dunrui
; APPLICANT: Drmanac, Radoje T.
; TITLE OF INVENTION: No. 6743619el Nucleic Acids and
; FILE REFERENCE: 802
; CURRENT APPLICATION NUMBER: US/09/774,528
; CURRENT FILING DATE: 2001-01-30
; NUMBER OF SEQ ID NOS: 441
; SOFTWARE: pt_FL_genes Version 2.0
; SEQ ID NO 366
; LENGTH: 2061
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (97)..(1206)
US-09-774-528-366

Query Match 74.8%; Score 17.2; DB 4; Length 2061;
Best Local Similarity 86.4%; Pred. No. 1.2e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCAACTACCACTG 22
|||||
Db 53 CCTTTTCGTCCTCACTTACACTG 32

FEATURE:
NAME/KEY: allele
LOCATION: 99117
OTHER INFORMATION: 5-130-276 : polymorphic base A or G
FEATURE:
NAME/KEY: allele
LOCATION: 103806
OTHER INFORMATION: 5-131-395 : polymorphic base A or T
FEATURE:
NAME/KEY: allele
LOCATION: 106940
OTHER INFORMATION: 5-133-375 : polymorphic base insertion of A
FEATURE:
NAME/KEY: allele
LOCATION: 108106
OTHER INFORMATION: 5-135-155 : polymorphic base insertion of A
FEATURE:
NAME/KEY: allele
LOCATION: 108149
OTHER INFORMATION: 5-135-198 : polymorphic base insertion of GTTT
FEATURE:
NAME/KEY: allele
LOCATION: 108308
OTHER INFORMATION: 5-135-357 : polymorphic base A or G
FEATURE:
NAME/KEY: allele
LOCATION: 108471
OTHER INFORMATION: 5-136-174 : polymorphic base C or T
FEATURE:
NAME/KEY: allele
LOCATION: 134134
OTHER INFORMATION: 5-140-120 : polymorphic base C or T
FEATURE:
NAME/KEY: allele
LOCATION: 134362
OTHER INFORMATION: 5-140-348 : polymorphic base insertion of A
FEATURE:
NAME/KEY: allele
LOCATION: 134374
OTHER INFORMATION: 5-140-361 : polymorphic base insertion of CA
FEATURE:
NAME/KEY: allele
LOCATION: 146328
OTHER INFORMATION: 5-143-84 : polymorphic base A or G
FEATURE:
NAME/KEY: allele
LOCATION: 146345
OTHER INFORMATION: 5-143-101 : polymorphic base A or C
FEATURE:
NAME/KEY: allele
LOCATION: 150329
OTHER INFORMATION: 5-145-24 : polymorphic base A or G
FEATURE:
NAME/KEY: allele
LOCATION: 160031
OTHER INFORMATION: 5-148-352 : polymorphic base G or T
FEATURE:
NAME/KEY: allele
LOCATION: 72771..72817
OTHER INFORMATION: polymorphic fragment 5-124-273 SEQ ID30
FEATURE:
NAME/KEY: allele
LOCATION: 72771..72817
OTHER INFORMATION: polymorphic fragment 5-124-273 SEQ ID51
FEATURE:
NAME/KEY: allele
LOCATION: 88050..88096
OTHER INFORMATION: polymorphic fragment 5-127-261 SEQ ID31
FEATURE:
NAME/KEY: allele
LOCATION: 88050..88096
OTHER INFORMATION: polymorphic fragment 5-127-261 SEQ ID52
FEATURE:

NAME/KEY: allele
LOCATION: 90819..90865
OTHER INFORMATION: complement polymorphic fragment 99-1437-325 SEQ ID49
FEATURE:
NAME/KEY: allele
LOCATION: 90819..90865
OTHER INFORMATION: complement polymorphic fragment 99-1437-325 SEQ ID70
FEATURE:
NAME/KEY: allele
LOCATION: 93690..93736
OTHER INFORMATION: polymorphic fragment 5-128-60 SEQ ID32
FEATURE:
NAME/KEY: allele
LOCATION: 93690..93736
OTHER INFORMATION: polymorphic fragment 5-128-60 SEQ ID53
FEATURE:
NAME/KEY: allele
LOCATION: 97099..97145
OTHER INFORMATION: polymorphic fragment 99-1442-224 SEQ ID50
FEATURE:
NAME/KEY: allele
LOCATION: 97099..97145
OTHER INFORMATION: polymorphic fragment 99-1442-224 SEQ ID71
FEATURE:
NAME/KEY: allele
LOCATION: 97130..97177
OTHER INFORMATION: polymorphic fragment 5-129-144 SEQ ID33
FEATURE:
NAME/KEY: allele
LOCATION: 97130..97177
OTHER INFORMATION: polymorphic fragment 5-129-144 SEQ ID54
FEATURE:
NAME/KEY: allele
LOCATION: 99075..99121
OTHER INFORMATION: polymorphic fragment 5-130-257 SEQ ID34
FEATURE:
NAME/KEY: allele
LOCATION: 99075..99121
OTHER INFORMATION: polymorphic fragment 5-130-257 SEQ ID55
FEATURE:
NAME/KEY: allele
LOCATION: 99094..99140
OTHER INFORMATION: polymorphic fragment 5-130-276 SEQ ID35
FEATURE:
NAME/KEY: allele
LOCATION: 99094..99140
OTHER INFORMATION: polymorphic fragment 5-130-276 SEQ ID56
FEATURE:
NAME/KEY: allele
LOCATION: 103783..103828
OTHER INFORMATION: polymorphic fragment 5-131-395 SEQ ID36
FEATURE:
NAME/KEY: allele
LOCATION: 103783..103828
OTHER INFORMATION: polymorphic fragment 5-131-395 SEQ ID57
FEATURE:
NAME/KEY: allele
LOCATION: 106918..106966
OTHER INFORMATION: polymorphic fragment 5-133-375 SEQ ID37
FEATURE:
NAME/KEY: allele
LOCATION: 106918..106966
OTHER INFORMATION: polymorphic fragment 5-133-375 SEQ ID58
FEATURE:
NAME/KEY: allele
LOCATION: 108084..108130
OTHER INFORMATION: polymorphic fragment 5-135-155 SEQ ID38
FEATURE:
NAME/KEY: allele
LOCATION: 108084..108130
OTHER INFORMATION: polymorphic fragment 5-135-155 SEQ ID59
FEATURE:
NAME/KEY: allele

```
; LOCATION: 108127..108177
; OTHER INFORMATION: polymorphic fragment 5-135-198 SEQ ID39
; FEATURE:
; NAME/KEY: allele
; LOCATION: 108127..108177
; OTHER INFORMATION: polymorphic fragment 5-135-198 SEQ ID60
; FEATURE:
```

```
Query Match          74.8%; Score 17.2; DB 3; Length 162450;
Best Local Similarity 86.4%; Pred. No. 2.2e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
```

```
QY      2 CTTTTCATCCAACTACCACTGA 23
Db      76273 CATTTCATCCAACTACCACTTA 76252
```

```
RESULT 13
US-09-949-016-14543/c
; Sequence 14543, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14543
; LENGTH: 387902
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(387902)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14543
```

```
Query Match          74.8%; Score 17.2; DB 4; Length 387902;
Best Local Similarity 86.4%; Pred. No. 2.5e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
```

```
QY      2 CTTTTCATCCAACTACCACTGA 23
Db      187998 CATTTCATCCAACTACCACTGA 187977
```

```
RESULT 14
US-09-949-016-12557/c
; Sequence 12557, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
```

```
; SOFTWARE: FastSeq for Windows Version 4.0
```

```
; SEQ ID NO 12557
```

```
; LENGTH: 421883
```

```
; TYPE: DNA
```

```
; ORGANISM: Human
```

```
; FEATURE:
```

```
; NAME/KEY: misc_feature
```

```
; LOCATION: (1)..(421883)
```

```
; OTHER INFORMATION: n = A,T,C or G
```

```
US-09-949-016-12557
```

```
Query Match          74.8%; Score 17.2; DB 4; Length 421883;
Best Local Similarity 86.4%; Pred. No. 2.5e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
```

```
QY      2 CTTTTCATCCAACTACCACTGA 23
Db      187998 CATTTCATCCAACTACCACTGA 187977
```

```
RESULT 15
US-09-328-352-3203/c
; Sequence 3203, Application US/09328352
; Patent No. 6562958
; GENERAL INFORMATION:
; APPLICANT: Gary L. Breton et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO ACINETOBACTER
; FILE REFERENCE: GTC99-03PA
; CURRENT APPLICATION NUMBER: US/09/328,352
; CURRENT FILING DATE: 1999-06-04
; NUMBER OF SEQ ID NOS: 8252
; SEQ ID NO 3203
; LENGTH: 942
; TYPE: DNA
; ORGANISM: Acinetobacter baumannii
US-09-328-352-3203
```

```
Query Match          73.0%; Score 16.8; DB 4; Length 942;
Best Local Similarity 90.0%; Pred. No. 1.7e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
```

```
QY      3 TTTTCATCCAACTACCACTG 22
Db      22 TTTTCATCGAATTACCACTG 3
```

```
Search completed: August 13, 2005, 06:48:49
Job time : 61.5213 secs
```

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 13, 2005, 03:26:33 ; Search time 1478.12 Seconds
(without alignments)
592.293 Million cell updates/sec

Title: US-10-673-854-2

Perfect score: 23

Sequence: 1 cctttcatcctaactaccactga 23

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: gb_est1:*

2: gb_est2:*

3: gb_hc3:*

4: gb_est3:*

5: gb_est4:*

6: gb_est5:*

7: gb_est6:*

8: gb_gss1:*

9: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	87.0	1018	8	BH163891 ENTTC11TF
2	19.4	84.3	592	1	AJ631826
3	18.8	81.7	404	2	BF048570 dc30f02.x
4	18.8	81.7	627	8	AZ334397 IM0083G14
5	18.8	81.7	644	6	CB102377 MWV_SQ008
6	18.8	81.7	860	9	CL072324 CH216-124
7	18.4	80.0	529	8	AZ926545 476_gis08
8	18.2	79.1	204	1	AA073416 mm84302.r
9	18.2	79.1	228	6	CD067162
10	18.2	79.1	281	2	BB486889
11	18.2	79.1	295	1	AV107656
12	18.2	79.1	394	8	BH170823
13	18.2	79.1	452	6	CB101123
14	18.2	79.1	581	9	CE706312 tigr-gss-
15	18.2	79.1	1184	8	CC320497
16	18.2	79.1	2145	3	AK044653
17	18	78.3	649	9	CG822989
18	17.8	77.4	279	3	CNS09JH9
19	17.8	77.4	404	8	AQ020753
20	17.8	77.4	410	3	CNS09JEW
21	17.8	77.4	422	8	AZ456776
22	17.8	77.4	497	8	B63439
23	17.8	77.4	500	8	AQ586167
24	17.8	77.4	528	1	AL926454

25	17.8	77.4	574	1	AV769356	AV769356 AV769356
26	17.8	77.4	576	4	BM646088	BM646088 170006873
27	17.8	77.4	577	7	CF447387	CF447387 EST683732
28	17.8	77.4	672	9	CR319820	CR319820 Medicago
29	17.8	77.4	722	6	CB818885	CB818885 EST 1775
30	17.8	77.4	752	9	AG376278	AG376278 Mus muscu
31	17.8	77.4	816	6	CB230720	CB230720 AGENCOURT
32	17.8	77.4	866	3	CNS09P2S	CBX068576 Single re
33	17.8	77.4	868	9	BX983278	BX983278 Forward s
34	17.8	77.4	877	3	CNS09P2R	BX068575 Single re
35	17.8	77.4	884	7	CR534985	CR534985 CR534985
36	17.8	77.4	931	3	CNS09P4Y	BX068870 Single re
37	17.8	77.4	947	3	CNS09S15	BX043209 Single re
38	17.8	77.4	954	3	CNS093GM	BX040562 Single re
39	17.8	77.4	954	8	CC208750	CC208750 CH261-101
40	17.8	77.4	985	9	AY412408	AY412408 Pan trogl
41	17.8	77.4	1022	3	CNS09PAX	BX068869 Single re
42	17.8	77.4	1053	3	CNS09J6C	BX060928 Single re
43	17.8	77.4	1122	4	BM563746	BM563746 AGENCOURT
44	17.4	75.7	220	9	CG590120	CG590120 OST243140
45	17.4	75.7	246	1	AV360610	AV360610 AV360610

ALIGNMENTS

RESULT 1
BH163891 1018 bp DNA linear GSS 24-SBP-2001
LOCUS ENTTC11TF Entamoeba histolytica Sheared DNA Entamoeba histolytica
DEFINITION genomic, genomic survey sequence.
ACCESSION BH163891
VERSION BH163891.1 GI:15737329
KEYWORDS GSS.
SOURCE Entamoeba histolytica
ORGANISM Entamoeba histolytica
REFERENCE 1 (bases 1 to 1018)
AUTHORS Loftus,B., Wang,Z., Van Aken,S. and Fraser,C.
TITLE Determination of clone end sequences from Entamoeba histolytica
JOURNAL HM1:IMSS sheared DNA library (2001)
COMMENT Contact: Brendan J Loftus
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0208
Fax: 301 838 3543
Email: bjloftus@tigr.org
Clones are derived from the Entamoeba histolytica HM1:IMSS sheared
DNA library
Seq primer: M13-Forward
Class: shotgun
High quality sequence start: 16
High quality sequence stop: 490.

FEATURES

source
1..1018
/organism="Entamoeba histolytica"
/mol_type="genomic DNA"
/strain="HM1:IMSS"
/db_xref="taxon:5759"
/clone_lib="Entamoeba histolytica Sheared DNA"
/note="Vector: pHOS1; Site 1: Bat 1; Constructed at The
Institute for Genomic Research (TIGR), Rockville, MD.
Genomic DNA isolated from broth cultures of E. histolytica
using a method described by Clark and Diamond (Clark,
C.G., and Diamond, L.S. (1993) Entamoeba histolytica: a
method for isolate identification. Exp. Parasitol.
77:450.). The DNA was mechanically sheared to give a
tight size distribution (~2 kb). The v + i method used for
the library construction is described in detail in Smith,
H.O. and Venter, J.C. (Making small insert libraries for
whole genome shotgun sequencing projects. In Genome

Sequencing: A Practical Approach, eds. M. Vaudin and B. Barrell, Oxford University Press, 1999).

ORIGIN
Query Match 87.0%; Score 20; DB 8; Length 1018;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCAACTACCAC 20
|||||
DB 777 CCTTTTCATCCAACTACCAC 796

RESULT 2
AJ631826/c 592 bp mRNA linear EST 21-APR-2004
LOCUS
DEFINITION
AJ631826 Prunus persica mesocarp S4 climacteric Prunus persica cDNA
clone Pp-S4EST0157, mRNA sequence.

ACCESSION
AJ631826
VERSION
AJ631826.1 GI:46472729

KEYWORDS
EST.

SOURCE
Prunus persica (peach)

ORGANISM
Prunus persica

Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids I; Rosales; Rosaceae; Amygdaloideae; Prunus.

REFERENCE
1 (bases 1 to 592)
Ziliotto, F., Begheldo, M., Rasori, A., Bonghi, C., Ramina, A. and
Tonutti, P.

TITLE
Microarray to study molecular and genetic aspects of ripening and
qualitative traits of peach (Prunus persica L. batesch) fruit

JOURNAL
Unpublished (2004)
CONTACT: Tonutti P
Environmental Agronomy and Crop Science
University of Padova
viale dell'Università, 16 Agripolis, Legnaro (PD) /35020, ITALY.

FEATURES
source

1..592
Location/Qualifiers
/organism="Prunus persica"
/mol_type="mRNA"
/cultivar="Pantasia"
/db_xref="taxon:3760"
/clone="Pp-S4EST0157"
/tissue_type="mesocarp"
/dev_stage="S4 climacteric"
/clone_lib="Prunus persica mesocarp S4 climacteric"

ORIGIN

Query Match 84.3%; Score 19.4; DB 1; Length 592;
Best Local Similarity 95.2%; Pred. No. 2.6e+02;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 CTTTTCATCCAACTACCAC 22
|||||
DB 419 CATTTCATCCAACTACCAC 399

RESULT 3
BF048570 404 bp mRNA linear EST 11-OCT-2000
LOCUS
DEFINITION
dc30f02.x1 NICHD_XGC_L11 Xenopus laevis cDNA clone IMAGE:3398619
3', mRNA sequence.

ACCESSION
BF048570
VERSION
BF048570.1 GI:10767073

KEYWORDS
EST.

SOURCE
Xenopus laevis (African clawed frog)

ORGANISM
Xenopus laevis

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Amphibia; Batrachia; Anura; Mesobatrachia; Pipidoidea; Pipidae;
Xenopodinae; Xenopus; Xenopus.

REFERENCE
1 (bases 1 to 404)
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

AUTHORS
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

JOURNAL
COMMENT

Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-re@mail.nih.gov
Tissue Procurement: Martha Rebert, Steven L. Klein, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: Xenopus clones from this library are available
through the I.M.A.G.E. Consortium/LLNL at: info@image.llnl.gov
Seq primer: -40UP from Gibco
High quality sequence stop: 403.

FEATURES
source

1..404
Location/Qualifiers
/organism="Xenopus laevis"
/mol_type="mRNA"
/db_xref="taxon:8355"
/clones="IMAGE:3398619"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NICHD_XGC_L11"
/note="Organ: liver; Vector: pCMV-SPORT6; Site_1: NotI;
Site 2: SalI; Cloned unidirectionally. Primer: Oligo dT.
Average insert size 1.4 kb. Constructed by Life
Technologies. Note: This is a Xenopus Gene Collection
(XGC) library."

ORIGIN

Query Match 81.7%; Score 18.8; DB 2; Length 404;
Best Local Similarity 90.9%; Pred. No. 4.8e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCAACTACCAC 22
|||||

DB 319 CTTTTCATCCAGTTACCAC 340
|||||

RESULT 4
AZ334397/c

LOCUS
DEFINITION
AZ334397 627 bp DNA linear GSS 29-SEP-2000
clone UUGC1M063G14 R, genomic survey sequence.

ACCESSION
AZ334397

VERSION
AZ334397.1 GI:10401685

KEYWORDS
GSS.

SOURCE
Mus musculus (house mouse)

ORGANISM
Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE
1 (bases 1 to 627)
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T.,
Reilly, M., Rose, R., Stokes, R., Tingey, A., von
Niederhausern, A. and Wright, D., Weiss, R.

Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center

RM. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA

Tel: 801 585 5606
Fax: 801 585 7177

Insert Length: 10000 Std Error: 0.00
Plate: 0063 row: G column: 14

Seq primer: CACACAGGAACAGCATGACC
Class: plasmid ends

High quality sequence stop: 627.
Location/Qualifiers
1..627

/organism="Mus musculus"
/mol_type="genomic DNA"

/strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0063G14"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /notes="Vector: FWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydronamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (G14732114|GB|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

ORIGIN

Query Match 81.7%; Score 18.8; DB 8; Length 627;
 Best Local Similarity 90.9%; Pred. No. 5.1e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 CTTTTCATCCAACTACCACTGA 23
 |||||
 Db 70 CTTTTCATCCAACTAGACTGA 49

RESULT 5

LOCUS CB102377 644 bp mRNA linear EST 28-JAN-2003
 DEFINITION MWV_SQ008044 AD-wrmcDNA Caenorhabditis elegans cDNA, mRNA sequence.
 ACCESSION CB102377
 VERSION CB102377.1 GI:27928184

KEYWORDS

SOURCE

ORGANISM

Caenorhabditis elegans
 Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida;
 Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
 1 (bases 1 to 644)

REFERENCE

AUTHORS

TITLE

Walhout,A.J., Sordella,R., Lu,X., Hartley,J.L., Temple,G.F.,
 Brasch,M.A., Thierry-Mieg,N. and Vidal,M.
 Protein interaction mapping in C. elegans using proteins involved
 in vulval development

JOURNAL

MEDLINE

PUBMED

COMMENT

Science 287 (5450), 116-122 (2000)
 20082953
 10615043
 Contact: Vidal M

Dana Farber Cancer Institute
 1 Jimmy Fund Way Smith 858, BOSTON, MA 02115, USA
 Tel: 617 632 5180
 Fax: 617 632 5739

Email: Marc.Vidal@dfci.harvard.edu
 Trace dvpl0604.x with Bait unknown
 POLYA=No.

FEATURES

source

Location/Qualifiers

1..644

/organism="Caenorhabditis elegans"

/mol_type="mRNA"

/strain="N2"

/db_xref="taxon:6239"

/sex="Hermaphrodite and male"

/tissue_type="whole animal"

/dev_stage="mixed stage"

/clone_lib="AD-wrmcDNA"
 /note="The AD-wrmcDNA library was generated with poly(A)+
 RNA isolated from both hermaphrodite and male N2 worms of
 all larval stages, embryos, adults and dauers and the
 subsequent generation of cDNAs by poly(A) priming. The
 cDNAs were cloned into pPC86"

ORIGIN

Query Match 81.7%; Score 18.8; DB 6; Length 644;
 Best Local Similarity 90.9%; Pred. No. 5.1e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 CTTTTCATCCAACTACCACTGA 23
 |||||
 Db 164 CTTTTCATCCAACTAGACTGA 185

RESULT 6

LOCUS

DEFINITION

CL072324 860 bp DNA linear GSS 31-DEC-2003
 CH216-124E11.RM1.1 CH216 Xenopus tropicalis genomic clone
 CH216-124E11 genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Xenopus tropicalis (western clawed frog)
 Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae;
 Xenopodinae; Xenopus; Silurana.

1 (bases 1 to 860)

Kremitzki,C., Carter,J., McPherson,J., Warren,W., Graves,T.,
 Mardis,E. and Wilson,R.

A physical map of the xenopus tropicalis genome

Unpublished (2003)

Contact: Richard K Wilson

Genome Sequencing Center

Washington University School of Medicine

Email: submissions@watson.wustl.edu

Insert Length: 175000 Std Error: 0.00

Seq primer: RM1 TAGCACTCACTATAGGAGA

Class: BAC ends

High quality sequence start: 102

High quality sequence stop: 445.

Location/Qualifiers

1..860

/organism="Xenopus tropicalis"

/mol_type="genomic DNA"

/strain="Nigerian frog"

/db_xref="taxon:8364"

/clone="CH216-124E11"

/sex="male"

/cell_line="Stock 248 F7A2, inbred N7"

/clone_lib="CH216"

/note="Vector: pTARBAC2.1; CHORI-216 Xenopus tropicalis
 BAC library"

ORIGIN

Query Match 81.7%; Score 18.8; DB 9; Length 860;
 Best Local Similarity 90.9%; Pred. No. 5.3e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCAACTACCACTG 22
 |||||
 Db 798 CCTTTCATCCAAATACCACTG 819

RESULT 7

LOCUS

DEFINITION

AZ926545 529 bp DNA linear GSS 01-APR-2001
 476.d1808d12.s1 Saccharomyces castellii NRRL Y-12630 Naumovia
 castellii genomic clone 476.d1808d12.s1, genomic survey sequence.

ACCESSION

```

VERSION  AZ926545.1  GI:13497447
KEYWORDS
SOURCE   Naumovia castellii
ORGANISM Naumovia castellii

REFERENCE
AUTHORS   Clifton,P.F., Hillier,L.W., Fulton,L., Graves,T., Miner,T.,
          Gish,W.R., Waterston,R.H. and Johnston,M.
TITLE     Surveying Saccharomyces genomes to identify functional elements by
          comparative DNA sequence analysis
JOURNAL   Unpublished (2001)
COMMENT   Contact: Johnston M
          Department of Genetics
          Washington University Medical School
          Box 8232, 4566 Scott Ave., St. Louis, MO 63110, USA
          Tel: 314 362 2735
          Fax: 314 362 7855
          Email: mj@genetics.wustl.edu
          Class: random plasmid subclone.
          Location/Qualifiers
            1..529
              /organism="Naumovia castellii"
              /mol_type="genomic DNA"
              /strain="NRRL Y-12630 (CBS 4309)"
              /db_xref="taxon:27288"
              /clone="476.d1808d12.81"
              /clone_lib="Saccharomyces castellii NRRL Y-12630"
              /note="Random genomic sequence"

ORIGIN
Query Match      80.0%; Score 18.4; DB 8; Length 529;
Best Local Similarity 95.0%; Pred. No. 7.7e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      4  TTTCATCCCAACTACCACTGA 23
        |||||
        |||||

DB      520  TTTCATCCCAAGTACCACTGA 501

RESULT 8
AA073416
LOCUS   mm84a02.r1 Stratagene mouse embryonic carcinomaRA (#937318) Mus
DEFINITION
ACCESSION AA073416
VERSION   AA073416.1  GI:1595145
KEYWORDS  EST.
SOURCE    Mus musculus (house mouse)
ORGANISM  Mus musculus

REFERENCE
AUTHORS   Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,
          Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,
          Schellenberg,K., Steptoe,M., Tan,P., Underwood,K., Moore,B.,
          Theising,B., Wyllie,T., Lennon,G., Soares,B., Wilson,R. and
          Waterston,R.
TITLE     The WashU-HMI Mouse EST Project
JOURNAL   Unpublished (1996)
COMMENT   Contact: Marra M/Mouse EST Project
          WashU-HMI Mouse EST Project
          Washington University School of MedicineP
          4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
          Tel: 314 286 1800
          Fax: 314 286 1810
          Email: mouseest@watson.wustl.edu
          This clone is available royalty-free through LLM; contact the
          IMAGE Consortium (info@image.llnl.gov) for further information.
          Seq primer: -28m13 rev1 ET from Amersham
          High quality sequence stop: 84.
          Location/Qualifiers

FEATURES
source
1..204
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="IMAGE:535082"
/tissue_type="carcinoma"
/dev_stage="embryonic"
/lab_host="SOUR (kanamycin resistant)"
/clone_lib="Stratagene mouse embryonic carcinomaRA
(#937318)"
/note="Vector: pBluescript SK-; Site 1: EcoRI; Site 2:
XhoI; Cloned unidirectionally. Primer: Oligo dt. p19 cell
line treated with retinoic acid. Average insert size: 1.0
kb; Uni-ZAP XR Vector; -5' adaptor sequence: 5'
GAATTCGGCAGGAG 3' -3' adaptor sequence: 5'
CTCGAGTTTTTTTTTTTTTTT 3'"

ORIGIN
Query Match      79.1%; Score 18.2; DB 1; Length 204;
Best Local Similarity 87.0%; Pred. No. 8.4e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY      1  CCTTTTCATCCCAACTACCACTGA 23
        |||||
        |||||

DB      110  CCTTTTCATCCCACTGCCTTTGA 132

RESULT 9
CD067162
LOCUS   MA1-0049T-R073-A09-U.G MA1-0049 Schistosoma mansoni CDNA clone
DEFINITION
ACCESSION MA1-0049T-R073-A09.G, mRNA sequence.
VERSION   CD067162
KEYWORDS  EST.
SOURCE    Schistosoma mansoni
ORGANISM  Schistosoma mansoni

REFERENCE
AUTHORS   Verjovski-Almeida,S., DeMarco,R., Martins,E.A.L., Guimaraes,P.E.M.,
          Ojopi,E.P.B., Paquola,A.C.M., Piazza,J.P., Nishiyama,M.Y. Jr.,
          Kitajima,J.P., Adamson,R.E., Ashton,P.D., Bonaldo,M.F.,
          Coulson,P.S., Dallon,G.P., Farias,L.P., Gregorio,S.P., Ho,P.L.,
          Leite,R.A., Malaquias,L.C.C., Marques,R.C.P., Miyasato,P.A.,
          Nascimento,A.L.T.O., Ohlweiler,F.P., Reis,E.M., Ribeiro,M.A.,
          Sa,R.G., Stukart,G.C., Soares,M.B., Gargioni,C., Kawano,T.,
          Rodrigues,V., Madeira,A.M.B.N., Wilson,R.A., Menck,C.P.M.,
          Setubal,J.C., Leite,L.C.C. and Dias-Neto,E.
TITLE     Transcriptome analysis of the acoelomate human parasite Schistosoma
          mansoni
JOURNAL   Nat. Genet. 35 (2), 148-157 (2003)
MEDLINE   22879926
PUBMED    12973350
COMMENT   Contact: Dr. Sergio Verjovski-Almeida
          Departamento de Bioquímica
          Instituto de Química - Universidade de Sao Paulo
          Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 Sao Paulo - SP,
          Brasil
          Tel: +55-11-3091-2173
          Fax: +55-11-3091-2186
          Email: verjo@iq.usp.br
          This sequence was derived from the FAPESP Schistosoma mansoni EST
          Genome Project. All sequences in the project were assembled and
          annotated. This entry and all the assembled sequences can be seen
          in the following URL http://bioinfo.iq.usp.br/schisto/
          Plate: MA1-0049T-R073 row: 9 column: A.
          Location/Qualifiers
            1..228
              /organism="Schistosoma mansoni"
              /mol_type="mRNA"
              /db_xref="taxon:6183"
              /clone="MA1-0049T-R073-A09.G"

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/db_xref="taxon:10090"
 /clone="2510040A06"
 /sex="mixed"
 /tissue_type="liver"
 /dev_stage="13-day embryo"
 /clone_lib="Mus musculus liver C57BL/6J 13-day embryo"

ORIGIN

Query Match 79.1%; Score 18.2; DB 1; Length 295;
 Best Local Similarity 87.0%; Pred. No. 8.9e+02;
 Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCCACTACCACTGA 23
 |||||
 Db 33 CCTTTTCATCCCACTGCCATGA 55

RESULT 12

BH170823
 LOCUS SALK_003398 Arabidopsis thaliana DNA linear GSS 03-OCT-2001
 DEFINITION thaliana genomic clone SALK_003398, genomic survey sequence.

ACCESSION BH170823
 VERSION BH170823.1 GI:15906526
 KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsids.

1 (bases 1 to 394)

AUTHORS Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
 Gadinab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
 Shinn,P., Zimmerman,J. and Ecker,J.R.

TITLE A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

JOURNAL

Unpublished (2001)

COMMENT

Contact: Joseph R. Ecker
 Salk Institute Genomic Analysis Laboratory (SIGNAL)
 The Salk Institute for Biological Studies
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
 Tel: 858 453 4100 x1752
 Fax: 858 558 6379
 Email: ecker@salk.edu
 This is single pass sequence recovered from the left border of
 TDNA.

Class: TDNA tagged.

Location/Qualifiers

1. .394
 /organism="Arabidopsis thaliana"
 /mol_type="genomic DNA"
 /ecotype="Col-0"
 /db_xref="taxon:3702"
 /clone="SALK_003398"

/note="PCR was performed on Arabidopsis thaliana lines
 each of which contains one or more TDNA insertion
 elements. The resultant fragment for each line was
 directly sequenced to determine the genomic sequence at
 the site of insertion. Details of the protocols used can
 be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN

Query Match 79.1%; Score 18.2; DB 8; Length 394;
 Best Local Similarity 87.0%; Pred. No. 9.2e+02;
 Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCCACTACCACTGA 23
 |||||
 Db 315 CATTTTATCCCACTACCACTAA 337

RESULT 13

CB101123/c
 LOCUS K16G09.y1 Ascaris suum embryo pAMP1 v2 Ascaris suum cDNA 5', mRNA
 DEFINITION sequence.

ACCESSION CB101123
 VERSION CB101123.1 GI:27926930
 KEYWORDS EST.

SOURCE Ascaris suum (pig roundworm)

ORGANISM

Eukaryota; Metazoa; Nematoda; Chromadorea; Ascaridida;

Ascaridoidea; Ascarididae; Ascaris.

1 (bases 1 to 452)

AUTHORS McCarter,J., Clifton,S., Chiapelli,B., Pape,D., Martin,J.,
 Wylie,T., Dante,M., Marra,M., Hillier,L., Kucaba,T., Theising,B.,
 Bowers,Y., Gibbons,M., Ritter,E., Bennett,J., Franklin,C.,
 Tsagarishvili,R., Ronko,I., Kennedy,S., Maguire,L., Beck,C.,
 Underwood,K., Steptoe,M., Allen,M., Person,B., Swaller,T.,
 Harvey,N., Schurk,R., Kohn,S., Shin,T., Jackson,Y., Cardenas,M.,
 McCann,R., Waterston,R. and Wilson,R.

TITLE The Washington Univ. Nematode EST Project, 1999
 JOURNAL Unpublished (1999)

COMMENT

Contact: McCarter JP

The Washington Univ. Nematode EST Project, 1999

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

The library was constructed by Claire Murphy and Dr. James McCarter
 at Washington University, St. Louis. The cDNA was made by using
 Dynabead oligo-dT priming (Dyna)l. PCR based library using a
 modified protocol from the SMART PCR cDNA Synthesis Kit from
 Clontech. Directionally cloned into the UDG sites of pAMP1. 30-60
 cell embryo material was provided by Dr. Richard Davis of City
 University of New York Graduate Center, College of Staten Island,
 Staten Island, NY (redavis@postbox.csi.cuny.edu).
 Seq primer: -40RP from Gibco

High quality sequence stop: 422.

FEATURES

source

Location/Qualifiers
 1. .452
 /organism="Ascaris suum"
 /mol_type="mRNA"
 /db_xref="taxon:6253"
 /dev_stage="30-60 cell embryo"
 /lab_host="DH10B Vector"
 /clone_lib="Ascaris suum embryo pAMP1 v2"
 /note="Vector: pAMP1; Site 1: NotI; Site 2: SalI; The
 library was constructed by Claire Murphy and Dr. James
 McCarter at Washington University, St. Louis. The cDNA was
 made by using Dynabead oligo-dT priming (Dyna)l. PCR based
 library using a modified protocol from the SMART PCR cDNA
 Synthesis Kit from Clontech. Directionally cloned into the
 UDG sites of pAMP1. 30-60 cell embryo material was
 provided by Dr. Richard Davis of City University of New
 York Graduate Center, College of Staten Island, Staten
 Island, NY (redavis@postbox.csi.cuny.edu).

ORIGIN

Query Match 79.1%; Score 18.2; DB 6; Length 452;
 Best Local Similarity 87.0%; Pred. No. 9.4e+02;
 Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CCTTTTCATCCCACTACCACTGA 23
 |||||
 Db 25 CCTTGTCATCGAATCATCACTGA 3

RESULT 14

CE706312/c

LOCUS

DEFINITION

tigr-gss-dog-17000369184273 Dog Library Canis familiaris genomic,

genomic survey sequence.

ACCESSION CE706312

```
VERSION CE706312.1 GI:37025703
KEYWORDS GSS.
SOURCE Canis familiaris (dog)
ORGANISM Canis familiaris
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
Kirkness, E.F., Bakna, V., Halpern, A.L., Levy, S., Remington, K.,
Rusch, D.B., Delcher, A.L., Pop, M., Wang, W., Fraser, C.M. and
Venter, J.C.
The dog genome: survey sequencing and comparative analysis
Science 301 (5641), 1898-1903 (2003)
MEDLINE 22875432
PUBMED 14512627
COMMENT Contact: Kirkness EF
The Institute for Genomic Research
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
Rockville, MD 20850, USA
Tel: 301-838-0200
Fax: 301-838-0208
Email: ekirknes@tigr.org
Class: shotgun.
FEATURES
    source
    Location/Qualifiers
    1..581
    /organism="Canis familiaris"
    /mol_type="genomic DNA"
    /strain="Standard Poodle"
    /db_xref="taxon:9615"
    /clone_lib="Dog Library"
    /note="Site 1: BstXI; Libraries were prepared from
    peripheral blood"
ORIGIN
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Best Local Similarity 87.0%; Pred. No. 9.7e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 CCTTTTCATCCCACTACCACTGA 23
Db 389 CCTTTTCATCCCACTACCACTGA 367
RESULT 15
LOCUS CC320497 1184 bp DNA linear GSS 14-MAY-2003
DEFINITION TAM32-27E3_Sp6.1 TAM32 Gallus gallus genomic clone TAM32-27E3,
genomic survey sequence.
ACCESSION CC320497
VERSION CC320497.1 GI:30714555
KEYWORDS GSS.
SOURCE Gallus gallus (chicken)
ORGANISM Gallus gallus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
Phasianinae; Gallus.
REFERENCE 1 (bases 1 to 1184)
AUTHORS Kremitzki, C., Higginbotham, J., Wylie, K., Carter, J., McPherson, J.,
Warren, W., Graves, T., Mardis, E. and Wilson, R.
TITLE Gallus gallus BAC End Reads
JOURNAL Unpublished (2003)
COMMENT Contact: Richard K. Wilson
Genome Sequencing Center
Washington University School of Medicine
Email: submissions@watson.wustl.edu
Insert Length: 152000 Std Error: 0.00
Seq primer: Sp6 ATTAGTGACACTATAG
Class: BAC ends
High quality sequence start: 8
High quality sequence stop: 751.
FEATURES
    source
    Location/Qualifiers
    1..1184
    /organism="Gallus gallus"
    /mol_type="genomic DNA"
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/strain="Red Jungle Fowl"
/db_xref="taxon:9031"
/clone="TAM32-27E3"
/sex="female"
/cell_line="UCD001, inbred 256"
/clone_lib="TAM32"
/note="Vector: pECBAC1; Site 1: EcoRI; Site 2: EcoRI;
TAM32 Female Chicken library - for library and clone
ordering information: http://www.hbz.tamu.edu"
ORIGIN
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Best Local Similarity 87.0%; Pred. No. 1.1e+03;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 CCTTTTCATCCCACTACCACTGA 23
Db 185 CATTCTCATCCCACTACCACTAA 207
Search completed: August 13, 2005, 06:44:57
Job time : 1486.12 secs
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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 13, 2005, 01:08:33 ; Search time 788.298 Seconds
(without alignments)
1536.704 Million cell updates/sec

Title: US-10-673-854-3

Perfect score: 25
Sequence: 1 tgaagaatttcagttcatagctgtg 25

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.*
1: gb_ba.*
2: gb_htg.*
3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_sta.*
12: gb_sy.*
13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	25	100.0	142099	9	AC002509
2	25	100.0	176872	9	AC012078
3	25	100.0	212280	9	HSB362E11
C 4	20.2	80.8	144552	8	CNS08CEK
C 5	20.2	80.8	155327	9	AC138761
C 6	20.2	80.8	157038	2	EX957355
C 7	20.2	80.8	167472	5	EX001041
C 8	20.2	80.8	198756	9	AC087575
C 9	20.2	80.8	199035	2	AC022263
C 10	20.2	80.8	204512	2	AC119029
C 11	20.2	80.8	215185	2	EX927073
C 12	20.2	80.8	226228	2	AC098027
C 13	19.8	79.2	44848	6	AR438843
C 14	19.8	79.2	57633	9	AC093382
C 15	19.8	79.2	72040	2	AC016347
C 16	19.8	79.2	81017	2	AC025010
C 17	19.8	79.2	91798	9	HSXB152G3
C 18	19.8	79.2	108418	9	AC007001
C 19	19.8	79.2	113587	9	AL590702

C 20	19.8	79.2	119024	9	AL159987	AL159987 Human DNA
C 21	19.8	79.2	124457	9	AC115115	AC115115 Homo sapi
C 22	19.8	79.2	137246	9	HS49J10	Z84572 Human DNA s
C 23	19.8	79.2	145859	9	AC009510	AC009510 Homo sapi
C 24	19.8	79.2	149188	9	AC114781	AC114781 Homo sapi
C 25	19.8	79.2	149403	2	AC130461	AC130461 Homo sapi
C 26	19.8	79.2	153468	9	AC113208	AC113208 Homo sapi
C 27	19.8	79.2	154439	2	AC073161	AC073161 Homo sapi
C 28	19.8	79.2	157267	9	AC008011	AC008011 Homo sapi
C 29	19.8	79.2	163777	9	AC067932	AC067932 Homo sapi
C 30	19.8	79.2	164858	9	AC026634	AC026634 Homo sapi
C 31	19.8	79.2	165558	4	AP006185	AP006185 Sus scrofa
C 32	19.8	79.2	167476	9	AC116038	AC116038 Homo sapi
C 33	19.8	79.2	167580	2	AC146124	AC146124 Pan trogl
C 34	19.8	79.2	168487	9	AL138820	AL138820 Human DNA
C 35	19.8	79.2	169416	9	AC022078	AC022078 Homo sapi
C 36	19.8	79.2	174365	2	AC024552	AC024552 Homo sapi
C 37	19.8	79.2	179755	9	AL603831	AL603831 Human DNA
C 38	19.8	79.2	180443	2	AC148366	AC148366 Callithri
C 39	19.8	79.2	184396	9	AC010853	AC010853 Homo sapi
C 40	19.8	79.2	184518	9	AC099524	AC099524 Homo sapi
C 41	19.8	79.2	188032	2	AC016393	AC016393 Homo sapi
C 42	19.8	79.2	189590	5	BX470167	BX470167 Zebrafish
C 43	19.8	79.2	198477	10	AC098390	AC098390 Rattus no
C 44	19.8	79.2	202950	9	AC017100	AC017100 Homo sapi
C 45	19.8	79.2	203270	9	AC064807	AC064807 Homo sapi

ALIGNMENTS

RESULT 1
AC002509
LOCUS
DEFINITION
AC002509
AC002509
VERSION
AC002509.1
KEYWORDS
HTG.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 142099)
AUTHORS
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
JOURNAL
Homo sapiens chromosome Y, clone 2Y
REFERENCE
2 (bases 1 to 142099)
AUTHORS
Hawkins,T.L., Birren,B.W., Faerman,K.H., Nusbaum,C., Lander,E.S., McKernan,K., Munro,C., Richardson,P., Barna,N., Chang,A., Cooke,P., Daly,M.J., Devon,K., Dewar,K., Forrest,C., Gage,D., Geraigery,K., Hagos,B., Huang,J., Hui,L., Jacotot,L., Kirby,A., Lane,M., MacKenzie,J., Marquis,N., McDermott,J., Molla,M., Morrow,J., Nachman,A., Naylor,J., Nusbaum,C., O'Connor,T., Olotu,A., Peterson,K., Reeve,M.P., Roberts,D., Rollins,G., Stillewell,J., Stone,C., Strickland,C., Sydney,K., Tang,L., Wilmer,F., Zemtseva,I., and Zody,M.
TITLE
Direct Submission
JOURNAL
Submitted (27-AUG-1997) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE
3 (bases 1 to 142099)
AUTHORS
Birren,B., Linton,L., Nusbaum,C., Allen,N., Anderson,M., Baker,J., Baldwin,J., Barna,N., Beckerly,R., Benn,J., Boutwell,C., Brown,A., Castle,A., Cerny,J., Colangelo,M., Collins,S., Collymore,A., Cooke,P., Corliss,D., Depayre,E., Devon,K., Dewar,K., Donelan,L., Ferreira,P., Fitzhugh,W., Forrest,C., Funke,R., Gage,D., Gardyna,S., Geraigery,K., Grant,G., Hagos,B., Heaford,A., Herena,L., Horton,L., Howland,J.C., Jacotot,L., Jones,C., Kann,L., Karatas,A., Lehoczy,J., Macdonald,P., Marquis,N., McSwan,P., McGurk,A., McKernan,K., Meldrim,J., Molla,M., Morris,W., Morrow,J., Mychaleckyj,J., Nahf,R., Naylor,J., Niloff,M., O'Connor,T., Roy,A., O'Donnell,P., Pavlin,B., Peterson,K., Riley,R., Roberts,D., Roy,A., Severy,P., Stange-Thomann,N., Stilwell,J., Stojanovic,N., Stone,C., Subramanian,A., Testaye,S., Tichovolsky,N., Torruella-Miller,I.,

Vassiliev,H., Vo.A., Wagner,A., Wheeler,J., Wu,Y., Wyman,D.,
 Ye,W.J., Zhao,J. and Zody,M.
 Direct Submission
 Submitted (25-NOV-1998) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 COMMENT
 On Nov 25, 1998 this sequence version replaced gi:3924665.
 All repeats were identified using RepeatMasker: Smit, A.F.A. &
 Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html.

FEATURES

Location/Qualifiers

source

1. .142099

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/map="Y"

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/clone_lib="unknown"

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complement(2160..2521)

/rpt_family="MER50"

2522..2578

/rpt_family="AluSg/x"

2859..2919

/rpt_family="MER51-internal"

complement(2920..2978)

/rpt_family="(CA)n"

2990..3699

/rpt_family="MER4-internal"

complement(3700..3814)

/rpt_family="LTR8"

3815..4468

/rpt_family="MER4-internal"

4523..7011

/rpt_family="MER4-internal"

complement(7295..7598)

/rpt_family="AluSp"

9162..9349

/rpt_family="MER4-internal"

9361..9509

/rpt_family="MER4A2"

9510..9811

/rpt_family="MER4A"

complement(9812..9962)

/rpt_family="L1M1"

complement(10212..10830)

/rpt_family="L1M4"

complement(10831..11120)

/rpt_family="AluY"

complement(11121..11337)

/rpt_family="L1M4"

complement(11508..12117)

/rpt_family="L1MEC"

complement(12315..12626)

/rpt_family="L1ME"

complement(12923..13136)

/rpt_family="MIR"

complement(13246..13328)

/note="Single-stranded terminator coverage."

13761..13782

/rpt_family="AT_rich"

14930..14950

/rpt_family="AT_rich"

15128..15231

/rpt_family="HAL1"

15292..15552

/rpt_family="HAL1"

complement(15690..16275)

/rpt_family="MER4B"

complement(16397..16551)

/rpt_family="MER66-internal"

16825..16846

repeat_region
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 repeat_region
 complement(17687..17920)
 /rpt_family="MER4-internal"
 repeat_region
 complement(18046..18789)
 /rpt_family="MER31-internal"
 repeat_region
 complement(19887..20110)
 /rpt_family="MER50"
 repeat_region
 complement(20133..20299)
 /rpt_family="MER4-internal"
 repeat_region
 20300..20593
 /rpt_family="AluY"
 repeat_region
 complement(20594..20673)
 /rpt_family="MER4-internal"
 repeat_region
 complement(20674..20974)
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 repeat_region
 complement(20975..21867)
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 repeat_region
 complement(21372..21498)
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 repeat_region
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 /rpt_family="MER4-internal"
 repeat_region
 complement(22252..22441)
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 repeat_region
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 /rpt_family="MER83-internal"
 repeat_region
 complement(22589..22693)
 /rpt_family="MER65-internal"
 repeat_region
 23324..23353
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 repeat_region
 complement(23437..23977)
 /rpt_family="MER4B"
 repeat_region
 complement(23978..24177)
 /rpt_family="MLT2A"
 repeat_region
 24557..24891
 /rpt_family="HAL1"
 repeat_region
 25476..25758
 /rpt_family="L2"
 repeat_region
 26069..26101
 /rpt_family="AT_rich"
 repeat_region
 26265..26489
 /rpt_family="MLTID"
 repeat_region
 26490..26517
 /rpt_family="(CAAA)n"
 repeat_region
 26518..26780
 /rpt_family="MLTID"
 repeat_region
 27156..27180
 /rpt_family="AT_rich"
 repeat_region
 complement(27417..27797)
 /rpt_family="L2"
 repeat_region
 complement(27963..28332)
 /rpt_family="MSTA"
 repeat_region
 complement(28333..29268)
 /rpt_family="MSTA-internal"
 repeat_region
 29269..29580
 /rpt_family="AluY"
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 complement(29581..30301)
 /rpt_family="MSTA-internal"
 repeat_region
 complement(30304..30679)
 /rpt_family="MSTA"
 repeat_region
 complement(30680..31159)
 /rpt_family="LTR40a"
 repeat_region
 complement(31325..31655)
 /rpt_family="L1ME"
 repeat_region
 31692..31731
 /rpt_family="AT_rich"
 repeat_region
 31883..31927
 /rpt_family="AT_rich"


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repeat_region complement(32107..33826)
repeat_region /rpt_family="L1M4"
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repeat_region 34987..35012
repeat_region /rpt_family="AT_rich"
repeat_region 35108..35669
repeat_region /rpt_family="MER4B"
repeat_region 36280..36322
repeat_region /rpt_family="AT_rich"
repeat_region complement(36385..36519)
repeat_region /rpt_family="MIR"
repeat_region complement(36755..37203)
repeat_region /rpt_family="MER4C"

Query Match 100.0%; Score 25; DB 9; Length 142099;
Best Local Similarity 100.0%; Pred. No. 1.7;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGAAGAAATTCAGTTCATAGCTTGT 25
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Db 35055 TGAAGAAATTCAGTTCATAGCTTGT 35079
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RESULT 2
AC012078 176872 bp DNA linear PRI 30-SEP-2000
LOCUS Homo sapiens BAC clone RP11-539022 from Y, complete sequence.
AC012078
ACCESSION AC012078.3 GI:7684580
VERSION
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 176872)
Sulston,J.E. and Waterston,R.
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
99063792
MEDLINE
PUBMED 9847074
REFERENCE 2 (bases 1 to 176872)
AUTHORS Hou,S., Maupin,R. and Gibson,A.
TITLE The sequence of Homo sapiens BAC clone RP11-539022
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 176872)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (19-OCT-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
4 (bases 1 to 176872)
Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (03-MAY-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
5 (bases 1 to 176872)
Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (12-JUN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
6 (bases 1 to 176872)
Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (30-SEP-2000) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On May 3, 2000 this sequence version replaced gi:7631053.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc

```

Contact: eapiens@watson.wustl.edu
 ----- Summary Statistics
 Center project name: H_NH0539022

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
 all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

The position of this clone was established as part of a collaboration between the Human Chromosome Y Mapping Project (Tomoko Kawaguchi, Helen Skaletsky, Laura G. Brown, Steve Rozen, and David C. Page at the Whitehead Institute for Biomedical Research, Cambridge MA) and the Washington University Genome Sequencing Center, St. Louis MO.

SOURCE INFORMATION:

The RP11-11 human BAC library was made from the blood of one male donor, as described by Osoegawa,K., Woon,P.Y., Zhao,B., Frengen,B., Tateno,M., Catanese,J.J. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (http://www.resgen.com) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (http://bacpac.med.buffalo.edu)

VECTOR: pSACE3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-530K5; the clone sequenced to the right is RP11-33605. Actual start of this clone is at base position 1 of RP11-539022; actual end is at base position 176872 of RP11-539022.

FEATURES	Location/Qualifiers
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	/db_xref="taxon:9606"
	/chromosome="Y"
	/map="Y"
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	/clone="RP11-539022"
	/clone_lib="RP11-11"
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repeat_region	202..1293
	/rpt_family="L1"
repeat_region	1294..1324
	/rpt_family=" (CAAA)n"
repeat_region	1325..2158
	/rpt_family="L1"
repeat_region	2159..2339
	/rpt_family="L1"
repeat_region	2340..3099
	/rpt_family="L1"
repeat_region	3120..3219
	/rpt_family="Alu"
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	/rpt_family="L1"
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	/rpt_family="L1"
repeat_region	4565..4619
	/rpt_family="Alu"
repeat_region	4638..5206
	/rpt_family="L1"
repeat_region	5255..5268

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120. .403
repeat_region /note="MRR39b repeat: matches 68. .453 of consensus"
404. .498
repeat_region /note="MRR39 repeat: matches 7. .103 of consensus"
502. .1573
repeat_region /note="LIM4 repeat: matches 4008. .5095 of consensus"
1569. .1914
repeat_region /note="LIM4 repeat: matches 3525. .3878 of consensus"
1906. .2066
repeat_region /note="LIMD repeat: matches 1376. .1528 of consensus"
2112. .2378
repeat_region /note="ALUJb repeat: matches 5. .274 of consensus"
2379. .2635
repeat_region /note="LIMD repeat: matches 1126. .1402 of consensus"
2651. .2697
repeat_region /note="MLT2D repeat: matches 504. .550 of consensus"
2840. .3252
repeat_region /note="MLT2D repeat: matches 77. .514 of consensus"
3351. .3420
repeat_region /note="MLT2D repeat: matches 1. .71 of consensus"
3421. .3902
repeat_region /note="HRRVL repeat: matches 5231. .5757 of consensus"
3903. .4462
repeat_region /note="280 copies 2 mer tt 57% conserved"
4465. .4782
repeat_region /note="ALUJo repeat: matches 1. .312 of consensus"
4783. .6107
repeat_region /note="HRRVL repeat: matches 3896. .5242 of consensus"
6238. .6378
repeat_region /note="LIMD repeat: matches 973. .1115 of consensus"
6640. .6783
repeat_region /note="LIMC5 repeat: matches 7751. .7910 of consensus"
7282. .7567
misc_feature /note="MLT1H repeat: matches 1. .302 of consensus"
complement (7766. .8138)
misc_feature /note="match: GSS: Em:AQ098528"
complement (7769. .8123)
misc_feature /note="match: GSS: Em:AQ105610"
complement (7779. .9205)
repeat_region /note="match: GSS: Em:AQ239326"
8002. .8469
repeat_region /note="MLT1D repeat: matches 1. .503 of consensus"
8477. .8654
repeat_region /note="MIR repeat: matches 75. .256 of consensus"
8715. .8812
gene /note="49 copies 2 mer aa 61% conserved"
complement (9391. .9893)
/genes="BB362B11.1"
CDS /pseudo
complement (9391. .9893)
/gene="BB362B11.1"
/note="BB362B11.1 (pseudogene similar to mouse GSG-154 and mosquito MRRG)
match: cDNAs: Em:AK001774
match: proteins: Sw:P50636 Tr:Q9Y4Y1 Tr:P90662 Tr:P91067"
/pseudo
/codon_start=1
/evidence=not_experimental
9919. .10251
misc_feature /note="match: STS: Em:T15642"
10912. .13011
repeat_region /note="L1 repeat: matches 2373. .4514 of consensus"
13018. .13758
repeat_region /note="TIGER1 repeat: matches 1. .774 of consensus"
13759. .13814
repeat_region /note="LIM4c repeat: matches 1557. .1609 of consensus"
13815. .14766
repeat_region /note="LITR5 repeat: matches 1. .969 of consensus"
14767. .14882
repeat_region /note="LIM4c repeat: matches 1609. .1737 of consensus"
15152. .15106
repeat_region /note="LIM3 repeat: matches 4670. .5642 of consensus"

repeat_region 16107. .16560
/note="LITR46 repeat: matches 1. .461 of consensus"
18749. .19851
repeat_region /note="HRRVH repeat: matches 3994. .5500 of consensus"
19925. .22648
repeat_region /note="HRRVH21 repeat: matches 1275. .3767 of consensus"
complement (22988. .23428)
misc_feature /note="match: GSS: Em:AQ619480"
23057. .23492
misc_feature /note="match: GSS: Em:AQ833119"
23749. .24192
repeat_region /note="LITR46 repeat: matches 1. .461 of consensus"
24194. .24348
repeat_region /note="LIM3 repeat: matches 5639. .5793 of consensus"
24386. .24485
repeat_region /note="LIM9 repeat: matches 5995. .6096 of consensus"
24497. .25002
repeat_region /note="LIM9 repeat: matches 5794. .6308 of consensus"
complement (25454. .26350)
misc_feature /note="match: GSS: Em:AQ894184"
25505. .25526
repeat_region /note="11 copies 2 mer ta 100% conserved"
25587. .25823
repeat_region /note="MIR repeat: matches 17. .262 of consensus"
complement (25882. .26240)
misc_feature /note="match: GSS: Em:AQ929805"
25885. .26180
misc_feature /note="match: GSS: Em:AQ754783"
complement (25910. .26219)
misc_feature /note="match: GSS: Em:AQ785533"
25945. .26256
misc_feature /note="match: GSS: Em:AQ628742"
25949. .26676
misc_feature /note="match: GSS: Em:ALI08886"
25950. .26147
misc_feature /note="match: GSS: Em:AQ230662"
25951. .26260
misc_feature /note="match: GSS: Em:AQ866229"
complement (25951. .26227)
misc_feature /note="match: GSS: Em:AQ541753"
complement (25951. .26105)
misc_feature /note="match: GSS: Em:AQ017857"
25951. .25978
repeat_region /note="14 copies 2 mer ta 92% conserved"
25954. .26352
misc_feature /note="match: GSS: Em:AQ525540"
complement (25978. .26215)
misc_feature /note="match: GSS: Em:AQ023352"
complement (25979. .26131)
misc_feature /note="match: GSS: Em:AQ322892"
25980. .26025
repeat_region /note="23 copies 2 mer ta 84% conserved"
complement (25984. .26583)
misc_feature /note="match: GSS: Em:AQ321139"
complement (25988. .26185)
misc_feature /note="match: GSS: Em:AQ226794"
25994. .26263
misc_feature /note="match: GSS: Em:AZ018667 Em:AZ052220"
complement (25998. .26169)
misc_feature /note="match: STS: Em:AU047022"
complement (26000. .26305)
misc_feature /note="match: GSS: Em:AQ123830"
26005. .26229
misc_feature /note="match: GSS: Em:AZ007683"
26009. .26160
misc_feature /note="match: STS: Em:HSB017WB9"
26009. .26147
misc_feature /note="match: GSS: Em:AZ069301"
26037. .26181
misc_feature /note="match: GSS: Em:AQ871738"
complement (26046. .26369)
misc_feature /note="match: GSS: Em:AQ827142"
26046. .26360

Query Match 100.0%; Score 25; DB 9; Length 212280;
 Best Local Similarity 100.0%; Pred. No. 1.6; 0; Indels 0; Gaps 0;
 Matches 25; Conservative 0; Mismatches 0;

QY 1 TGAAGAAATTCAGTTCATAGCTTGT 25
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 Db 116062 TGAAGAAATTCAGTTCATAGCTTGT 116086
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RESULT 4
 CNS08CBK/c
 LOCUS
 DEFINITION Oryza sativa chromosome 12, linear PLN 21-NOV-2003
 from chromosome 12 of cultivar Nipponbare of ssp. japonica of Oryza
 sativa (rice), complete sequence.
 AL928747
 AL928747.3 GI:28412540

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

Oryza sativa (japonica cultivar-group)
 Oryza sativa (japonica cultivar-group)
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Ehrhartoideae; Oryzaceae; Oryza.

REFERENCE
 1 (bases 1 to 144552)
 Choisne,N., Orjeda,G., Cattolico,L., Demange,N., Wincker,P.,
 Seguren,B., Pelletier,E., Scarpelli,C., Salanoubat,M.,
 Weissenbach,J. and Quetier,F.

Oryza sativa chromosome 12 sequencing

Unpublished

2 (bases 1 to 144552)

Genoscope.

Direct Submission

Submitted (20-NOV-2003) Genoscope - Centre National de Sequencage :
 BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
 - Web : www.genoscope.cns.fr)

On Feb 18, 2003 this sequence version replaced gi:24430303.

Center: Genoscope / Centre National de Sequencage

Center code: GS

Web site: http://www.genoscope.cns.fr/

Contact: SeqRef@genoscope.cns.fr

The following sequence is oriented from the T7 to the SP6 end. The
 nucleotide sequence of this BAC clone was generated by combining
 Syngenta and Genoscope sequencing data.
 Upstream BAC (overlapping the T7 end) : P0185F07 (AC=BX667525)
 Downstream BAC (overlapping the SP6 end) : OJ1111_F12 (AC=AL732536)

 FINISHING BOUNDARIES
 FINISHED SEGMENT STARTS AT BASE 1
 FINISHED SEGMENT ENDS AT BASE 110216

FEATURES
 source

Location/Qualifiers
 1. .144552
 /organism="Oryza sativa (japonica cultivar-group)"
 /mol_type="genomic DNA"
 /cultivar="Nipponbare"
 /sub_species="japonica"
 /db_xref="taxon:39947"
 /chromosome="12"
 /clone="OSJNBa0017A21"
 /clone_lib="OSJNBa"

ORIGIN

Query Match 80.8%; Score 20.2; DB 8; Length 144552;
 Best Local Similarity 88.0%; Pred. No. 2e+02;
 Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTCATAGCTTGT 25
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 Db 66142 TGAAGAAATTCAGTTCATAGCTTGT 66118
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RESULT 5.

AC138761/c
 LOCUS
 DEFINITION
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

AC138761 155327 bp DNA linear PRI 01-MAR-2003
 Homo sapiens chromosome 17, clone RP11-1109M24, complete sequence.
 AC138761
 HTG.
 AC138761.4 GI:28626680
 Homo sapiens (human)

REFERENCE
 AUTHORS
 TITLE
 JOURNAL
 REFERENCE
 AUTHORS

1 (bases 1 to 155327)
 Birren,B., Nusbaum,C. and Lander,E.

Homo sapiens chromosome 17, clone RP11-1109M24

Unpublished

2 (bases 1 to 155327)

Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
 Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B.,
 Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collamore,A.,
 Cook,A., Cooke,P., DeArelano,K., Dewar,K., Diaz,J.S., Dodge,S.,
 Faros,S., Ferreira,P., FitzGerald,M., Gage,D., Galagan,J.,
 Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hafez,N.,
 Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
 Kamat,A., Karatas,A., Kellis,C., Landers,T., Levine,R.,
 Lindblad-Toh,K., Liu,G., MacLean,C., Macdonald,P., Major,J.,
 Matthews,C., McCarthy,M., Meldrim,J., Meneus,L., Mihova,T.,
 Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C.,
 Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J.,
 Peterson,K., Phunkhang,P., Pierre,N., Raymond,C., Retta,R.,
 Rise,C., Rogov,P., Roman,J., Roy,A., Schauer,S., Schuback,R.,
 Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N.,
 Stojanovic,N., Talamas,J., Teefaye,S., Theodore,J., Topham,K.,
 Travers,M., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X.,
 Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission

Submitted (16-JAN-2003) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA

3 (bases 1 to 155327)

Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N.,
 Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T.,
 Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y.,
 Collamore,A., Cook,A., Cooke,P., Corum,B., DeArelano,K.,
 Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faros,S.,
 Ferreira,P., FitzGerald,M., Gage,D., Galagan,J., Gardyna,S.,
 Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B.,
 Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
 Kamat,A., Karatas,A., Kellis,C., Landers,T., Levine,R.,
 Lindblad-Toh,K., Liu,G., Lui,A., Mabbitt,R., MacLean,C.,
 Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M.,
 Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J.,
 Nguyen,C., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P.,
 O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
 Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P.,
 Roman,J., Schauer,S., Schuback,R., Seaman,S., Severy,P., Smith,C.,
 Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M.,
 Talamas,J., Teefaye,S., Theodore,J., Topham,K., Travers,M.,
 Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X.,
 Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission

Submitted (13-FEB-2003) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA

4 (bases 1 to 155327)

Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N.,
 Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T.,
 Boguslavskiy,L., Boukhgalter,B., Camarata,J., Chang,J., Choepel,Y.,
 Collamore,A., Cook,A., Cooke,P., Corum,B., DeArelano,K.,
 Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faros,S.,
 Ferreira,P., FitzGerald,M., Gage,D., Galagan,J., Gardyna,S.,
 Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B.,
 Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
 Kamat,A., Karatas,A., Kellis,C., Landers,T., Levine,R.,
 Lindblad-Toh,K., Liu,G., Lui,A., Mabbitt,R., MacLean,C.,
 Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M.,
 Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J.,

Nguyen,C., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P.,
 O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
 Rachupka,A., Ramaamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P.,
 Roman,J., Schauer,S., Schupback,R., Seaman,S., Severy,P., Smith,C.,
 Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M.,
 Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M.,
 Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X.,
 Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
 Direct Submission
 Submitted (01-MAR-2003) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Mar 1, 2003 this sequence version replaced gi:28301978.
 All repeats were identified using RepeatMasker:
 Smith, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu
 ----- Project Information
 Center project name: L29291
 Center clone name: 1109_M_24

FEATURES

Location/Qualifiers

Source
 1. .155327
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosome="17"
 /map="17"
 /clone="RP11-1109M24"
 /clone_lib="RPC1-11 Human Male BAC"
 17..72
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 72..145
 /rpt_family="(TA)n"
 complement(291..500)
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 527..568
 /rpt_family="AT-rich"
 3043..3079
 /rpt_family="U2"
 3307..3333
 /rpt_family="(TTTTTG)n"
 3543..3849
 /rpt_family="AluY"
 4155..4182
 /rpt_family="AT-rich"
 4757..4780
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 5591..5653
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 complement(7305..7355)
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 complement(7485..8409)
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 9060..9088
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 10340..10370
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 complement(10371..10522)
 /rpt_family="AluSg/x"
 complement(10831..11204)
 /rpt_family="THE1C"
 complement(11205..12793)
 /rpt_family="THE1C-int"
 complement(12794..13177)
 /rpt_family="THE1C"
 13179..13980
 /rpt_family="L2"

repeat_region 14702..15028
 /rpt_family="MER2"
 repeat_region 15087..15562
 /rpt_family="MER70A"
 repeat_region 15761..15933
 /rpt_family="L1ME3B"
 repeat_region 15967..16422
 /rpt_family="L1ME3B"
 repeat_region 16436..16459
 /rpt_family="AT-rich"
 repeat_region 16712..17467
 /rpt_family="L1ME3B"
 complement(17468..17947)
 /rpt_family="MLT1D"
 repeat_region 17948..18034
 /rpt_family="L1ME3B"
 repeat_region 18747..18836
 /rpt_family="AT-rich"
 complement(19087..19362)
 /rpt_family="AluX"
 repeat_region complement(19585..19878)
 /rpt_family="AluY"
 repeat_region complement(19890..20046)
 /rpt_family="AluSp"
 repeat_region complement(22092..22373)
 /rpt_family="AluX"
 repeat_region 23331..23384
 /rpt_family="AT-rich"
 complement(23597..23644)
 /rpt_family="LTR66"
 repeat_region 24249..24549
 /rpt_family="AluSp"
 repeat_region 24778..24804
 /rpt_family="(CA)n"
 repeat_region 24855..25198
 /rpt_family="THE1B"
 repeat_region 25208..26788
 /rpt_family="THE1B-int"
 repeat_region 26790..26993
 /rpt_family="THE1B"
 repeat_region 27626..27661
 /rpt_family="AT-rich"
 repeat_region 28903..28931
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 complement(29301..29597)
 /rpt_family="AluSc"
 repeat_region complement(29687..29914)
 /rpt_family="L1MA3"
 repeat_region 30315..30352
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 repeat_region 30632..30682
 /rpt_family="AT-rich"
 complement(30713..31593)
 /rpt_family="L1MA3"
 repeat_region 31594..32226

Query Match 80.8%; Score 20.2; DB 9; Length 155327;

Best Local Similarity 88.0%; Pred.No.2e+02; 3; Indels 0; Gaps 0;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTCATAGCTTGT 25

Db 72347 TGAAGAAATTCAGTTCATAGCTTGT 72323

RESULT 6

BX957355

LOCUS

DEFINITION

unordered pieces.

ACCESSION

VERSION

KEYWORDS

BX957355 157038 bp DNA linear HTG 01-MAR-2004
 Danio rerio clone CH211-66115, *** SEQUENCING IN PROGRESS ***
 HTG; HTGS_PHASE1.

```

SOURCE
ORGANISM      Danio rerio (zebrafish)
               Danio rerio
               Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
               Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
               Cypriniformes; Cyprinidae; Danio.
REFERENCE      1 (bases 1 to 157038)
AUTHORS        McLay, K.
TITLE          Direct Submission
JOURNAL        Submitted (26-FEB-2004) Wellcome Trust Sanger Institute, Hinxton,
               Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
               zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
COMMENT        On Mar 1, 2004 this sequence version replaced gi:42821059.
               ----- Genome Center
               Center: Wellcome Trust Sanger Institute
               Center code: SC
               Web site: http://www.sanger.ac.uk
               Contact: zfish-help@sanger.ac.uk
               ----- Project Information
               Center project name: ZC66115
               ----- Summary Statistics
               Assembly program: XGAP4; version 4.5
               Chemistry: Dye-terminator; 100% of reads
               Consensus quality: 153845 bases at least Q40
               Consensus quality: 154522 bases at least Q30
               Consensus quality: 155205 bases at least Q20
               Insert size: 156338; sum-of-contigs
               Insert size: 175167; 1.9% error; agarose-fp
               Quality coverage: 10.89x in Q20 bases; sum-of-contigs Quality
               coverage: 9.88x in Q20 bases; agarose-fp
               -----
               * NOTE: This is a 'working draft' sequence. It currently
               * consists of 8 contigs. The true order of the pieces
               * is not known and their order in this sequence record is
               * arbitrary. Gaps between the contigs are represented as
               * runs of N, but the exact sizes of the gaps are unknown.
               * This record will be updated with the finished sequence
               * as soon as it is available and the accession number will
               * be preserved.
               *
               1 58492: contig of 58492 bp in length
               * 58493 58592: gap of 100 bp
               * 77232 77232: contig of 18640 bp in length
               * 77233 77332: gap of 100 bp
               * 77333 86310: contig of 8978 bp in length
               * 86311 86410: gap of 100 bp
               * 86411 103070: contig of 16660 bp in length
               * 103071 103170: gap of 100 bp
               * 103171 134661: contig of 31491 bp in length
               * 134662 134761: gap of 100 bp
               * 134762 142097: contig of 7336 bp in length
               * 142098 142197: gap of 100 bp
               * 142198 150392: contig of 8195 bp in length
               * 150393 150492: gap of 100 bp
               * 150493 157038: contig of 6546 bp in length.
               Location/Qualifiers
               1..157038
               /organism="Danio rerio"
               /mol_type="genomic DNA"
               /db_xref="taxon:7955"
               /clone="CH211-66115"
               /clone_lib="CHORI-211"
               1..58492
               /note="assembly_fragment:01643
               clone_end:SP6
               vector_side:left"
               58593..77232
               /note="assembly_fragment:00719
               fragment_chain:1"
               77233..86310
               /note="assembly_fragment:00337
               fragment_chain:1"
               86411..103070
               /note="assembly_fragment:00475"
               103171..134661

```

```

/note="assembly_fragment:01114
fragment_chain:2"
134762..142097
/note="assembly_fragment:00031
fragment_chain:2"
142198..150392
/note="assembly_fragment:00222
fragment_chain:2"
150493..157038
/note="assembly_fragment:00124
fragment_chain:2
clone_end:T7
vector_side:right"

ORIGIN

Query Match      80.8%; Score 20.2; DB 2; Length 157038;
Best Local Similarity 88.0%; Pred. No. 2e+02;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGAGAAATTCAGTTCATAGCTTGT 25
   ||| ||||| ||||| ||||| |||||
Db 40806 TGAGGAAATTCAGTTTATAGCTTTT 40830

RESULT 7
BX001041/c
LOCUS BX001041 167472 bp DNA linear VRT 13-MAR-2003
DEFINITION Zebrafish DNA sequence from clone CH211-62M7, complete sequence.
ACCESSION BX001041
VERSION BX001041.6 GI:28971483
KEYWORDS HTG.
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
          Cypriniformes; Cyprinidae; Danio.
REFERENCE 1 (bases 1 to 167472)
AUTHORS Pelan,S.
TITLE Direct Submission
JOURNAL Submitted (13-MAR-2003) Wellcome Trust Sanger Institute, Hinxton,
          Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
          zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
          On Mar 14, 2003 this sequence version replaced gi:28552286.
COMMENT ----- Genome Center
          Center: Wellcome Trust Sanger Institute
          Center code: SC
          Web site: http://www.sanger.ac.uk
          Contact: zfish-help@sanger.ac.uk
          -----
          During sequence assembly data is compared from overlapping clones.
          Where differences are found these are annotated as variations
          together with a note of the overlapping clone name. Note that the
          variation annotation may not be found in the sequence submission
          corresponding to the overlapping clone, as we submit sequences with
          only a small overlap as described above.
          This sequence was finished as follows unless otherwise noted: all
          regions were either double-stranded or sequenced with an alternate
          chemistry or covered by high quality data (i.e., phred quality >=
          30); an attempt was made to resolve all sequencing problems, such
          as compressions and repeats; all regions were covered by at least
          one plasmid subclone or more than one M13 subclone; and the
          assembly was confirmed by restriction digest, except on the rare
          occasion of the clone being a YAC.
          The following abbreviations are used to associate primary accession
          numbers given in the feature table with their source databases:
          Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information
          on the WORMPEP database can be found at
          http://www.sanger.ac.uk/Projects/Celegans/wormpep Repeat names
          beginning 'Dr' were identified by the Recon repeat discovery system
          (Zhirong Bao and Sean Eddy, submitted), and those beginning 'drr'
          were identified by Rick Waterman (Stephen Johnson lab, WashU). For
          further information see http://www/Projects/D_rerio/fishmask.shtml
          CH211-62M7 is from a CHORI-211 BAC library

```


repeat_region	/rpt_family="AluSq" complement(5197..5481)	repeat_region	19930..20237 /rpt_family="AluSq"
repeat_region	/rpt_family="AluSx" complement(5707..5838)	Query Match	80.8%; Score 20.2; DB 9; Length 198756;
repeat_region	/rpt_family="FLAM_C" 6298..6597	Best Local Similarity	88.0%; Pred. No. 1.9e+02;
repeat_region	/rpt_family="AluJb" 6600..7279	Matches	22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
repeat_region	/rpt_family="Tigger1" complement(7280..7570)	QY	1 TGAAGAAATTCAGTTCATGCTTGT 25
repeat_region	/rpt_family="AluSp" 7628..7821	Db	143321 TGAAGAAATTCAGTTCATGGCTTTT 143345
repeat_region	/rpt_family="MER115" 8003..8103	RESULT 9	
repeat_region	/rpt_family="(TTATA)n" 8125..8626	AC022263	
repeat_region	/rpt_family="Tigger1" complement(8627..8904)	LOCUS	
repeat_region	/rpt_family="AluSx" 8905..9037	DEFINITION	AC022263 Homo sapiens chromosome UL clone RP11-305L6, WORKING DRAFT
repeat_region	/rpt_family="Tigger1" complement(9041..9331)	ACCESSION	AC022263.4 GI:7630939
repeat_region	/rpt_family="LTR16C" 9518..9606	VERSION	HTG; HTGS_PHASE1; HTGS_DRAFT.
repeat_region	/rpt_family="MER115" 9723..10065	KEYWORDS	Homo sapiens (human)
repeat_region	/rpt_family="MER115" complement(10467..10750)	SOURCE	Homo sapiens
repeat_region	/rpt_family="LIMA4" complement(11038..11252)	ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
repeat_region	/rpt_family="MIR" 11357..11671	REFERENCE	1 (bases 1 to 199035) Waterston,R.H.
repeat_region	/rpt_family="AluSx" complement(11766..11866)	AUTHORS	The sequence of Homo sapiens clone
repeat_region	/rpt_family="L2" complement(12415..12629)	TITLE	Unpublished
repeat_region	/rpt_family="MIR" 12775..12805	JOURNAL	2 (bases 1 to 199035) Waterston,R.H.
repeat_region	/rpt_family="AT_rich" 13003..13046	AUTHORS	Direct Submission
repeat_region	/rpt_family="AT_rich" complement(13289..13376)	TITLE	Submitted (27-JAN-2000) Genome Sequencing Center, Washington
repeat_region	/rpt_family="LIPAL6" 13617..13754	JOURNAL	University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
repeat_region	/rpt_family="FLAM_A" complement(13756..14094)	COMMENT	On Apr 21, 2000 this sequence version replaced gi:7230974.
repeat_region	/rpt_family="MER93" 14538..14558		
repeat_region	/rpt_family="AT_rich" 14571..15059		
repeat_region	/rpt_family="LIMB3" 15060..15383		
repeat_region	/rpt_family="AluSg" 15384..15959		
repeat_region	/rpt_family="LIMB3" 15968..16030		
repeat_region	/rpt_family="AT_rich" 16512..16538		
repeat_region	/rpt_family="AT_rich" complement(16541..16839)		
repeat_region	/rpt_family="AluY" 17404..17700		
repeat_region	/rpt_family="AluSg" 17717..17993		
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repeat_region	/rpt_family="(TAAAA)n" complement(18048..18322)		
repeat_region	/rpt_family="LIMD1" complement(18493..18674)		
repeat_region	/rpt_family="AluSg/x" 18800..19089		
repeat_region	/rpt_family="AluJb"		
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----- Genome Center -----			
Center: Washington University Genome Sequencing Center			
Center code: WUGSC			
Web site: http://genome.wustl.edu/gsc/index.shtml			
----- Project Information -----			
Center project name: H NH0305L06			
----- Summary Statistics -----			
Sequencing vector: M13; 61%			
Sequencing vector: plasmid; 39%			
Chemistry: Dye-primer ET; 61% of reads			
Chemistry: Dye-terminator Big Dye; 39% of reads			
Assembly program: Phrap; version 0.990319			
Consensus quality: 197548 bases at least Q40			
Consensus quality: 197985 bases at least Q30			
Consensus quality: 198133 bases at least Q20			
Insert size: 211000; agarose-fp			
Insert size: 198435; sum-of-contigs			
Quality coverage: 6.81 in Q20 bases; agarose-fp			
Quality coverage: 7.25 in Q20 bases; sum-of-contigs			

* NOTE: This is a 'working draft' sequence. It currently			
* consists of 7 contigs. The true order of the pieces			
* is not known and their order in this sequence record is			
* arbitrary. Gaps between the contigs are represented as			
* runs of N, but the exact sizes of the gaps are unknown.			
* This record will be updated with the finished sequence.			
* as soon as it is available and the accession number will			
* be preserved.			
* 1 3010: contig of 3010 bp in length			
* 3011 3110: gap of unknown length			
* 3111 15619: contig of 12509 bp in length			
* 15620 15719: gap of unknown length			
* 15720 27681: contig of 11962 bp in length			
* 27682 27781: gap of unknown length			
* 27782 50877: contig of 23096 bp in length			
* 50878 50977: gap of unknown length			
* 50978 83000: contig of 32023 bp in length			


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* 83001 83100: gap of unknown length
* 83101 118000: contig of 34900 bp in length
* 118001 118100: gap of unknown length
* 118101 199035: contig of 80935 bp in length.
FEATURES
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                /db_xref="taxon:9606"
                /chromosome="UL"
                /clone="RP11-305L6"
ORIGIN
Query Match      80.8%; Score 20.2; DB 2; Length 199035;
Best Local Similarity 88.0%; Pred. No. 1.9e+02;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTCATAGCTTGT 25
Db 30469 TGAAGAAATTCAGTTCATAGCTTTT 30493

RESULT 10
AC119029/c
LOCUS AC119029 204512 bp DNA linear HTG 20-NOV-2002
DEFINITION Rattus norvegicus clone CH230-50113, *** SEQUENCING IN PROGRESS
***
AC119029
VERSION AC119029.4 GI:25137856
KEYWORDS HTG; HTGS_Phas22; HTGS_DRAFT; HTGS_ENRICHED.
SOURCE Rattus norvegicus (Norway rat)
ORGANISM Rattus
            Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
            Rattus.
REFERENCE 1 (bases 1 to 204512)
AUTHORS Muzny,D.,Marie., Metzker,M.,Lee., Abramson,S., Adams,C., Alder,J.,
        Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
        Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
        Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
        Biswal,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
        Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
        Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
        Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,
        Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
        Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
        Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
        Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
        Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
        Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
        Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
        Gebregeorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
        Gunaratne,P., Haaland,W., Hamil,C., Hamilton,C., Hamilton,K.,
        Harvey,Y., Havlak,P., Haves,A., Henderson,N., Hernandez,J.,
        Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hogues,M.,
        Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A.,
        Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
        Karpach,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
        Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
        Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
        Lorensuhewa,L., Louised,H., Lozado,R.J., Lu,X., Ma,J.,
        Maheshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A.,
        Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
        Mawhney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
        Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
        Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L.,
        Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,
        Nwaokelimeh,O., Okwuonu,G., Olarnpunsagoon,A., Pal,S., Parks,K.,
        Pasternak,S., Paul,H., Perez,A., Perez,L., Pfannkuch,C.,
        Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L.-L.,
        Fuzao,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
        Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
        Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,

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TITLE JOURNAL
REFERENCE 2 (bases 1 to 204512)
AUTHORS Worley,K.C.
TITLE JOURNAL
SUBMITTED (24-APR-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 204512)
Rat Genome Sequencing Consortium.
Direct Submission
SUBMITTED (20-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Nov 20, 2002 this sequence version replaced gi:23908182.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GVZK
Center clone name: CH230-50113
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 166862 bases at least Q40
Consensus quality: 170683 bases at least Q30
Consensus quality: 173177 bases at least Q20
Estimated insert size: 174509; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 204512: contig of 204512 bp in length.
Location/Qualifiers
1..204512
/mol_type="genomic DNA"
/db_xref="taxon:10113"
/clone="CH230-50113"
FEATURES
    source

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Sanders,W., Savary,G., Scherer,S., Scott,G., Shatsman,S., Shen,H., Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D., Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Soosa,J., Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C., Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K., Valas,R., Vera,V., Villaseana,D., Waldron,L., Walker,B., Wang,J., Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F., Williams,G., Willson,R., Wlaczek,R., Wooden,H., Worley,K., Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V., Yu,F., Zhang,J., Zhou,X., Zhao,S., Zhao,S., Dunn,D., von Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O., Weinstock,G. and Gibbs,R.A.

Direct Submission
Unpublished
2 (bases 1 to 204512)
Worley,K.C.

Direct Submission
Submitted (24-APR-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 204512)
Rat Genome Sequencing Consortium.

Direct Submission
Submitted (20-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Nov 20, 2002 this sequence version replaced gi:23908182.

The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GVZK
Center clone name: CH230-50113
----- Summary Statistics

Assembly program: Phrap; version 0.990329
Consensus quality: 166862 bases at least Q40
Consensus quality: 170683 bases at least Q30
Consensus quality: 173177 bases at least Q20
Estimated insert size: 174509; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 204512: contig of 204512 bp in length.

Location/Qualifiers
1..204512
/mol_type="genomic DNA"
/db_xref="taxon:10113"
/clone="CH230-50113"

```

misc_feature 1. 1705
/note="wgs_end_extension
clone_end:Sp6"
misc_feature 3869. 4734
/note="clone_boundary
clone_end:Sp6
site:
end_sequence:B2277134"
64608. 66128
/note="wgs_contig"
misc_feature 166948. 168569
/note="wgs_contig"
misc_feature 168900. 170461
/note="wgs_contig"
misc_feature 192738. 193528
/note="clone_boundary
clone_end:T7
site:
end_sequence:B2277133"
196728. 198414
/note="wgs_end_extension
clone_end:T7"
misc_feature 199983. 201685
/note="wgs_end_extension
clone_end:T7"
misc_feature 201736. 202797
/note="wgs_end_extension
clone_end:T7"
misc_feature 203408. 204512
/note="wgs_end_extension
clone_end:T7"

ORIGIN
Query Match 80.8%; Score 20.2; DB 2; Length 204512;
Best Local Similarity 88.0%; Pred. No. 1.9e+02;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 TGAGAAATTCAGTTTCATAGCTTGT 25
|||||
Db 198373 TGAGAAATTCATTTTCATAGGTGT 198349

RESULT 11
BX927073 215185 bp DNA linear HTG 18-FEB-2004
LOCUS
DEFINITION
Danio rerio clone DKEYP-78C2, *** SEQUENCING IN PROGRESS ***, 7
unordered pieces.
ACCESSION BX927073.3 GI:42592534
VERSION
KEYWORDS HTG; HTGS PHASE1.
SOURCE
Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.
REFERENCE
1 (bases 1 to 215185)
McLay, K.
Direct Submission
Submitted (16-FEB-2004) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Feb 17, 2004 this sequence version replaced gi:41016204.
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: zfish-help@sanger.ac.uk
----- Project Information
Center project name: zKp78C2
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Chemistry: Dye-terminator; 100% of reads
Consensus quality: 211838 bases at least Q40

misc_feature 1. 1705
/note="wgs_end_extension
clone_end:Sp6"
misc_feature 3869. 4734
/note="clone_boundary
clone_end:Sp6
site:
end_sequence:B2277134"
64608. 66128
/note="wgs_contig"
misc_feature 166948. 168569
/note="wgs_contig"
misc_feature 168900. 170461
/note="wgs_contig"
misc_feature 192738. 193528
/note="clone_boundary
clone_end:T7
site:
end_sequence:B2277133"
196728. 198414
/note="wgs_end_extension
clone_end:T7"
misc_feature 199983. 201685
/note="wgs_end_extension
clone_end:T7"
misc_feature 201736. 202797
/note="wgs_end_extension
clone_end:T7"
misc_feature 203408. 204512
/note="wgs_end_extension
clone_end:T7"

ORIGIN
Query Match 80.8%; Score 20.2; DB 2; Length 204512;
Best Local Similarity 88.0%; Pred. No. 1.9e+02;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 TGAGAAATTCAGTTTCATAGCTTGT 25
|||||
Db 198373 TGAGAAATTCATTTTCATAGGTGT 198349

RESULT 11
BX927073 215185 bp DNA linear HTG 18-FEB-2004
LOCUS
DEFINITION
Danio rerio clone DKEYP-78C2, *** SEQUENCING IN PROGRESS ***, 7
unordered pieces.
ACCESSION BX927073.3 GI:42592534
VERSION
KEYWORDS HTG; HTGS PHASE1.
SOURCE
Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.
REFERENCE
1 (bases 1 to 215185)
McLay, K.
Direct Submission
Submitted (16-FEB-2004) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Feb 17, 2004 this sequence version replaced gi:41016204.
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: zfish-help@sanger.ac.uk
----- Project Information
Center project name: zKp78C2
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Chemistry: Dye-terminator; 100% of reads
Consensus quality: 211838 bases at least Q40

```

Consensus quality: 212487 bases at least Q30
 Consensus quality: 213129 bases at least Q20
 Insert size: 214585; sum-of-contigs
 Insert size: 219324; 2.6% error; agarose-fp
 Quality coverage: 8.00x in Q20 bases; sum-of-contigs Quality
 coverage: 7.92x in Q20 bases; agarose-fp

 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 7 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence.
 * as soon as it is available and the accession number will
 * be preserved.

```

* 1 3729: contig of 3729 bp in length
* 3730 3829: gap of 100 bp
* 3830 26383: contig of 22554 bp in length
* 26384 26483: gap of 100 bp
* 26484 33119: contig of 6636 bp in length
* 33120 33219: gap of 100 bp
* 33220 110818: contig of 77599 bp in length
* 110819 127149: gap of 100 bp
* 127150 127249: contig of 16231 bp in length
* 127250 154645: contig of 27396 bp in length
* 154646 154745: gap of 100 bp
* 154746 215185: contig of 60440 bp in length.

```

FEATURES

```

Location/Qualifiers
1..215185
/organism="Danio rerio"
/mol_type="genomic DNA"
/db_xref="taxon:7955"
/clone="DKEYP-78C2"
/clone_lib="DanioKeyPilot"
1..3729
/note="assembly_fragment:00023
fragment_chain:1"
3830..26383
/note="assembly_fragment:00331
fragment_chain:1"
26484..33119
/note="assembly_fragment:00064
fragment_chain:2"
33220..110818
/note="assembly_fragment:01713
fragment_chain:2"
110319..127149
/note="assembly_fragment:00146"
127250..154645
/note="assembly_fragment:00622.0"
154746..215185
/note="assembly_fragment:00949"

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ORIGIN

Query Match 80.8%; Score 20.2; DB 2; Length 215185;
 Best Local Similarity 88.0%; Pred. No. 1.9e+02;
 Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

```

Qy 1 TGAGAAATTCAGTTTCATAGCTTGT 25
|||||
Db 124798 TGAGAAATTCAGTTTATAGCTTTT 124822

```

RESULT 12

```

AC098027
LOCUS
DEFINITION
Rattus norvegicus clone CH230-53E3, *** SEQUENCING IN PROGRESS ***,
2 unordered pieces.
ACCESSION
AC098027
VERSION
HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.
KEYWORDS
Rattus norvegicus (Norway rat)
SOURCE

```

ORGANISM

Rattus norvegicus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus

REFERENCE
AUTHORS

1 (bases 1 to 226228)
Muzny, D. Marie, Metzker, M. Lee, Abramson, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,
Blawie, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,
Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,
Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,
Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,
Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,
Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,
Egan, A., Escotto, M., Eugene, C., Evans, C. A., Falls, T., Fan, G.,
Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,
Fraser, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M.,
Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W.,
Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K.,
Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J.,
Hernandez, R., Hines, S., Hladun, S. L., Hodgson, A., Hogues, M.,
Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A.,
Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,
Karpach, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C.,
Kowis, C., Kraft, C. L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J.,
Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,
Lorensueta, L., Louised, H., Lozada, R. J., Lu, X., Ma, J.,
Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A.,
Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E.,
Mawhney, S., McLeod, M. P., McNeill, T. Z., Meenen, E.,
Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,
Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L.,
Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S.,
Nwaokeme, O., Okwuon, G., Olarnpungsoon, A., Pal, S., Parks, K.,
Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C.,
Plopper, F., Polinder, A., Popovic, D., Primus, E., Pu, L. L.,
Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M. A., Reigh, R.,
Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,
Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S. J.,
Sanders, W., Savary, G., Scherer, S., Scott, G., Shatsman, S., Shen, H.,
Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C. D., Smaj, D.,
Sneed, A., Sodergren, E., Song, X. Z., Sorelle, R., Soes, J.,
Steinle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C.,
Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmari, K.,
Valas, R., Vera, V., Villaseana, D., Waldron, L., Walker, B., Wang, J.,
Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, P.,
Williams, G., Willson, R., Wleczky, R., Wooden, H., Worley, K.,
Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,
Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von
Niederhausern, A., Weiss, R., Smith, D. R., Holt, R. A., Smith, H. O.,
Weinstock, G. and Gibbs, R. A.

TITLE
JOURNAL

Unpublished
Direct Submission
2 (bases 1 to 226228)
Worley, K. C.

REFERENCE
AUTHORS

Submitted (23-OCT-2001) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 226228)

REFERENCE
AUTHORS

Rat Genome Sequencing Consortium.
Direct Submission

TITLE
JOURNAL

Submitted (10-MAY-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

COMMENT

On May 10, 2003 this sequence version replaced gi:23096288.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled with Atlas
(<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described
in the feature table below represents a scaffold in the Atlas

assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GFYN
Center clone name: CH230-53E3
----- Summary Statistics
Assembly program: Atlas 3.0;
Consensus quality: 217218 bases at least Q40
Consensus quality: 219723 bases at least Q30
Consensus quality: 221029 bases at least Q20
Estimated insert size: 228733; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 2 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 224885: contig of 224885 bp in length
* 224986 224985: gap of unknown length
* 224986 226228: contig of 1243 bp in length.

FEATURES
source

Location/Qualifiers
1..226228
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-53E3"
1..1471
/note="wgs contig"
4273..5861
/note="wgs_contig"

misc_feature

misc_feature

ORIGIN

Query Match 80.8%; Score 20.2; DB 2; Length 226228;
Best Local Similarity 88.0%; Pred. No. 1.9e+02;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTTCATAGCTGT 25
|||||
Db 183838 TGAAGAAATTCATTCATAGAGTGT 183862
|||||

RESULT 13

AR438843/c
LOCUS
DEFINITION
Sequence 42 from patent US 6664105.
AR438843
ACCESSION
AR438843.1 GI:42663846
VERSION
KEYWORDS
Unknown.
SOURCE
ORGANISM
Unknown.

AR438843 44848 bp DNA linear PAT 20-FEB-2004
Sequence 42 from patent US 6664105.

AR438843 1 GI:42663846
KEYWORDS
Unknown.
SOURCE
ORGANISM
Unknown.

REFERENCE
1 (bases 1 to 44848)
Pecker, I., Vlodavsky, I. and Feinstein, E.
Polynucleotide encoding a polypeptide having heparanase activity
and expression of same in genetically modified cells
Patent: US 6664105-A 42 16-DEC-2003;
JOURNAL

```

FEATURES                               Location/Qualifiers
Source                                  1..44848
/mol_type="genomic DNA"

ORIGIN
Query Match                          79.2%; Score 19.8; DB 6; Length 44848;
Best Local Similarity                91.3%; Pred. No. 3.5e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||
Db 30803 GAAGAAATTCAGTTCATGTTG 30781

RESULT 14
AC093382 LOCUS                               57633 bp DNA linear PRI 10-JAN-2002
DEFINITION Homo sapiens BAC clone RP11-476D14 from 2, complete sequence.
AC093382 AC093382
VERSION AC093382.3 GI:17978444
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
AUTHORS Sulston,J.E. and Waterston,R.
TITLE Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE 99063792
PUBMED 9847074
REFERENCE
AUTHORS Radlonsen,M., Kozlowski,A. and Doebber,A.
TITLE The sequence of Homo sapiens BAC clone RP11-476D14
JOURNAL Unpublished (2002)
REFERENCE
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (21-AUG-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (26-DEC-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (03-JAN-2002) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (10-JAN-2002) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
COMMENT On Dec 26, 2001 this sequence version replaced gi:17227294.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@watson.wustl.edu
----- Summary Statistics
Center project name: H_NH0476D14
-----
NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
clone sections once, or longer because we provide a small overlap
between neighboring data submissions.

```

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO.. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RPC11 human BAC library was made from the blood of one male donor, as described by Osoegawa,K., Woon,P.Y., Zhao,B., Frengen,E., Tateno,M., Catanese,J.J. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the right is RP11-504L12, 2000 bp overlap finished in RP11-476D14. Actual start of this clone is at base position 1 of RP11-476D14; actual end is at base position 106332 of RP11-504L12.

FEATURES

	Location/Qualifiers
source	1..57633
	/organism="Homo sapiens"
	/mol_type="genomic DNA"
	/db_xref="taxon:9606"
	/chromosome="2"
	/map="2"
	/clone="RP11-476D14"
	/clone_lib="RPC11"
	217..580
repeat_region	/rpt_family="L2"
repeat_region	581..879
repeat_region	/rpt_family="Alu"
repeat_region	880..1165
repeat_region	/rpt_family="L2"
repeat_region	1279..1413
repeat_region	/rpt_family="MIR"
repeat_region	1572..2071
repeat_region	/rpt_family="MaLR"
repeat_region	2133..2163
repeat_region	/rpt_family="(TTTC)n"
repeat_region	2137..2447
repeat_region	/rpt_family="Alu"
repeat_region	3163..3423
repeat_region	/rpt_family="Alu"
repeat_region	3418..3450
repeat_region	/rpt_family="AT-rich"
repeat_region	3437..3712
repeat_region	/rpt_family="L1"
repeat_region	3547..3588
repeat_region	/rpt_family="(TG)n"
repeat_region	3923..3945
repeat_region	/rpt_family="AT-rich"
repeat_region	4058..4291
repeat_region	/rpt_family="MER1_type"
repeat_region	4309..4553
repeat_region	/rpt_family="L1"
repeat_region	4575..5148
repeat_region	/rpt_family="L1"
repeat_region	5136..5173
repeat_region	/rpt_family="AT-rich"
repeat_region	6280..6376
repeat_region	/rpt_family="MER1_type"

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/rpt_family="MIR"
repeat_region 6922. .7276
/rpt_family="MaLR"
repeat_region 8012. .8033
/rpt_family="AT_rich"
repeat_region 8015. .8308
/rpt_family="Alu"
repeat_region 8900. .9227
/rpt_family="L2"
repeat_region 11593. .11747
/rpt_family="CRI"
repeat_region 13142. .13247
/rpt_family="MIR"
repeat_region 13782. .13898
/rpt_family="MIR"
repeat_region 15262. .15515
/rpt_family="MIR"
repeat_region 15577. .15642
/rpt_family="MER1_type?"
repeat_region 15647. .15837
/rpt_family="MIR"
misc_feature 15974. .16439
/notes="similar to Mus musculus EST BB614050
(NID:G16454511)"
repeat_region 17442. .18075
/rpt_family="ERV1"
repeat_region 19208. .19494
/rpt_family="ERV1"
repeat_region 19495. .20184
/rpt_family="ERV1"
repeat_region 20185. .20438
/rpt_family="ERV1"
repeat_region 20449. .20592
/rpt_family="L1"
repeat_region 20932. .21491
/rpt_family="ERV1"
repeat_region 21492. .21533
/rpt_family="AT_rich"
repeat_region 23942. .24226
/rpt_family="Alu"
repeat_region 24369. .24735
/rpt_family="ERV1"
repeat_region 25188. .25292
/rpt_family="MaLR"
repeat_region 25293. .25606
/rpt_family="Alu"
repeat_region 25450. .25481
/rpt_family="(T)n"
misc_feature 25457. .25472
/notes="match to EST BG167210 (NID:G12673913)"
repeat_region 25607. .25924
/rpt_family="MaLR"
repeat_region 26111. .26503
/rpt_family="ERV1"
repeat_region 26667. .27022
/rpt_family="ERV1"
repeat_region 27061. .27121
/rpt_family="ERV1"
repeat_region 27131. .27355
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repeat_region 27651. .27720
/rpt_family="Alu"
repeat_region 27795. .27847
/rpt_family="L2"
repeat_region 28053. .28276
/rpt_family="Alu"
repeat_region 29103. .29243
/rpt_family="(GAAA)n"
repeat_region 29284. .29501
/rpt_family="MER1_type"

```

Query Match

79.2%; Score 19.8; DB 9; Length 57633;

```

Best Local Similarity 91.3%; Pred. No. 3.4e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTTCATGCTTG 24
    |||||
Db 19627 GAAGAAATTCAGTTTCATGCTTG 19649

RESULT 15
AC016347
LOCUS AC016347
DEFINITION Homo sapiens chromosome 15 clone RP11-189L10 map 15, LOW-PASS
SEQUENCE SAMPLING.
AC016347
AC016347.2 GI:9141889
VERSION HTG; HTGS PHASE0.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
1 (bases 1 to 72040)
Homo sapiens chromosome 15, clone RP11-189L10
Unpublished
2 (bases 1 to 72040)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
Baldwin,J., Barna,N., Beckerly,R., Boguslavskiy,L., Boukhgalter,B.,
Brown,A., Castle,A., Collangelo,M., Collins,S., Collymore,A.,
Cooke,P., DeArellano,K., Dewar,K., Domino,M., Donegan,L., Doyle,M.,
Ferreira,P., Fitzhugh,W., Forrest,C., Funke,R., Gage,D., Hagan,M.,
Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karats,A., Klein,J.,
Lehoczky,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N.,
McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J.,
Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
Peterson,K., Pollara,V., Riley,K., Roy,A., Santos,R., Severy,P.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tesfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.
Direct Submission
Submitted (24-NOV-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jul 13, 2000 this sequence version replaced gi:6466986.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L4952
Center clone name: 189_L_10
-----
* NOTE: This record contains 81 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
* 1 807: contig of 807 bp in length
* 808 907: gap of 100 bp
* 908 1700: contig of 793 bp in length
* 1701 1800: gap of 100 bp
* 1801 2587: contig of 787 bp in length
* 2588 2687: gap of 100 bp
* 2688 3476: contig of 789 bp in length

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* 3477 3576: gap of 100 bp
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* 5210 5309: gap of 100 bp
* 5310 6079: contig of 770 bp in length
* 6080 6179: gap of 100 bp
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* 8753 8852: gap of 100 bp
* 8853 9634: contig of 782 bp in length
* 9635 9734: gap of 100 bp
* 9735 10519: contig of 785 bp in length
* 10520 10619: gap of 100 bp
* 10620 11415: contig of 796 bp in length
* 11416 11515: gap of 100 bp
* 11516 12314: contig of 799 bp in length
* 12315 12414: gap of 100 bp
* 12415 13201: contig of 787 bp in length
* 13202 13301: gap of 100 bp
* 13302 14092: contig of 791 bp in length
* 14093 14192: gap of 100 bp
* 14193 14981: contig of 789 bp in length
* 14982 15081: gap of 100 bp
* 15082 15849: contig of 768 bp in length
* 15850 15949: gap of 100 bp
* 15950 16775: contig of 826 bp in length
* 16776 16875: gap of 100 bp
* 16876 17667: contig of 792 bp in length
* 17668 17767: gap of 100 bp
* 17768 18574: contig of 807 bp in length
* 18575 18674: gap of 100 bp
* 18675 19478: contig of 804 bp in length
* 19479 19578: gap of 100 bp
* 19579 20380: contig of 802 bp in length
* 20381 20480: gap of 100 bp
* 20481 21287: contig of 807 bp in length
* 21288 21387: gap of 100 bp
* 21388 22175: contig of 788 bp in length
* 22176 22275: gap of 100 bp
* 22276 23029: contig of 754 bp in length
* 23030 23129: gap of 100 bp
* 23130 23914: contig of 785 bp in length
* 23915 24014: gap of 100 bp
* 24015 24788: contig of 774 bp in length
* 24789 24888: gap of 100 bp
* 24889 25686: contig of 798 bp in length
* 25687 25786: gap of 100 bp
* 25787 26593: contig of 807 bp in length
* 26594 27521: contig of 828 bp in length
* 27522 27621: gap of 100 bp
* 27622 28381: contig of 767 bp in length
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* 30294 31069: contig of 776 bp in length
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* 35669 36472: contig of 804 bp in length
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* 36573 37360: contig of 788 bp in length
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* 46385 47191: contig of 807 bp in length
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* 49055 49838: contig of 784 bp in length
* 49839 49938: gap of 100 bp
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* 50828 51634: contig of 807 bp in length
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* 51735 52530: contig of 796 bp in length
* 52531 52630: gap of 100 bp
* 52631 53412: contig of 782 bp in length
* 53413 53512: gap of 100 bp
* 53513 54297: contig of 785 bp in length
* 54298 54397: gap of 100 bp
* 54398 55184: contig of 787 bp in length
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* 55285 56093: contig of 809 bp in length
* 56094 56193: gap of 100 bp
* 56194 57001: contig of 808 bp in length
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* 64100 64199: gap of 100 bp

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Beat Local Similarity 91.3%; Pred. No. 3.3e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||
Db 71709 GAAGAAATTCAGTTCATAGCTTG 71731

Search completed: August 13, 2005, 05:04:24
Job time : 797.298 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 12, 2005, 22:21:57 ; Search time 200.266 Seconds
(without alignments)
738.985 Million cell updates/sec

Title: US-10-673-854-3

Perfect score: 25

Sequence: 1 tgaagaattcagttcatagctgtg 25

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 segs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N Geneseq_16Dec04:*

- 1: Geneseqn1980s:*
- 2: Geneseqn1990s:*
- 3: Geneseqn2000s:*
- 4: Geneseqn2001as:*
- 5: Geneseqn2001bs:*
- 6: Geneseqn2002as:*
- 7: Geneseqn2002bs:*
- 8: Geneseqn2003as:*
- 9: Geneseqn2003bs:*
- 10: Geneseqn2003cs:*
- 11: Geneseqn2003ds:*
- 12: Geneseqn2004as:*
- 13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	19.8	79.2	922	12	ADJ43067
C 2	19.8	79.2	44848	3	AAA75080
C 3	19.8	79.2	44848	10	ADG88832
C 4	19.8	79.2	44848	12	ADL16411
C 5	19.8	79.2	44848	12	ADM48748
C 6	19.8	79.2	183178	10	ADL13873
C 7	19.2	76.8	49507	11	ACN44572
C 8	19.2	76.8	202001	6	ABS52506
C 9	19.2	76.8	202001	10	ADG46742
C 10	19.2	76.0	235	12	ADP92027
C 11	18.8	75.2	1227	11	ACH97547
C 12	18.8	75.2	176771	12	ADQ97167
C 13	18.6	74.4	4212	6	ABZ14865
C 14	18.6	74.4	116858	11	ACN44212
C 15	18.2	72.8	978	10	ADK54409
C 16	18.2	72.8	5536	4	AAK89473
C 17	18.2	72.8	40463	11	ACN44904
C 18	18.2	72.8	98638	12	ADQ97919
C 19	18.2	72.8	164991	10	ADL13635
C 20	18.2	72.8	176080	12	ADL08124

C 21	18	72.0	520	5	AAS84456
C 22	18	72.0	2509	12	ADQ63965
C 23	17.8	71.2	491	4	ABA57693
C 24	17.8	71.2	491	4	AAI37269
C 25	17.8	71.2	491	4	AAK31375
C 26	17.8	71.2	491	4	AAK05751
C 27	17.8	71.2	491	4	ABS31056
C 28	17.8	71.2	491	6	ABS06128
C 29	17.8	71.2	708	13	ADS60583
C 30	17.8	71.2	1325	4	AAH52803
C 31	17.8	71.2	1485	6	ABN91876
C 32	17.8	71.2	1485	13	ADS01312
C 33	17.8	71.2	2978	4	AAH54735
C 34	17.8	71.2	3745	4	AAH54681
C 35	17.6	70.4	384	8	ABX50480
C 36	17.6	70.4	462	6	ABN26076
C 37	17.6	70.4	515	12	ADJ10653
C 38	17.6	70.4	545	13	ADQ50923
C 39	17.6	70.4	1259	6	ABA01916
C 40	17.6	70.4	1999	10	ADC08531
C 41	17.6	70.4	3089	13	ADS54895
C 42	17.6	70.4	3270	13	ACN38754
C 43	17.6	70.4	3808	5	ADL62420
C 44	17.6	70.4	90220	6	ABK83576
C 45	17.6	70.4	245531	13	ABD33022

ALIGNMENTS

RESULT 1

ADJ43067/c
ID ADJ43067 standard; cDNA; 922 BP.

AC ADJ43067;

DT 06-MAY-2004 (first entry)

DE Plant cDNA #4067.

KW Plant; gene; ss; transcription; plant genome augmentation; cereal;
KW soybean; alfalfa; sunflower; canola; cotton; peanut; tobacco; sugar beet;
KW maize; barley; sorghum; rice; wheat; crop plant; insecticide resistance;
KW stress tolerance; salt tolerance; cold tolerance; drought tolerance;
KW plant nutrition; apical dominance; dwarfism; early flowering; antiviral;
KW antifungal.

OS Eukaryota.

XX US2004016025-A1.

XX 22-JAN-2004.

XX 26-SEP-2002; 2002US-00260238.

XX 26-SEP-2001; 2001US-0325277P.

XX 26-SEP-2001; 2001US-0325448P.

XX 04-APR-2002; 2002US-0370620P.

XX (BUDW/) BUDWORTH P.

XX (MOUG/) MOUGHAMER T.

XX (BRIG/) BRIGGS S P.

XX (COOP/) COOPER B.

XX (GLAZ/) GLAZERBOOK J.

XX (GOFF/) GOFF S A.

XX (KATA/) KATAGIRI F.

XX (KREP/) KREPS J.

XX (PROV/) PROVART N.

XX (RICK/) RICHE D.

XX (ZHUT/) ZHU T.

Budworth P, Moughamer T, Briggs SP, Cooper B, Glazebrook J;
PI Goff SA, Katagiri F, Kreps J, Provart N, Ricke D, Zhu T;

XX WPI; 2004-190374/18.

XX

XX New rice promoter, useful for manipulating crop plants to alter or

PT improve phenotypic characteristics, e.g. produce large quantities of oil

PT or proteins, resistance to insecticides, virus or fungi, stress tolerance

PT or high nutritional value.

XX

XX Example 13; SEQ ID NO 4067; 230pp; English.

XX

XX The invention relates to plant nucleotide sequences that direct seed-,

CC leaf- and/or stem-, panicle-, root- or pollen-specific or -preferential

CC or constitutive transcription of an operatively linked nucleic acid

CC segment. The invention also relates to a method for augmenting a plant

CC genome and a method of identifying a gene, where its expression is

CC altered in the seed, leaf, stem, panicle, pollen, root or is constitutive

CC in a plant cell. The plant is a cereal, e.g. soybean, alfalfa, sunflower,

CC canola, cotton, peanut, tobacco or sugar beet, preferably maize, barley,

CC sorghum, rice or wheat. The polynucleotides and the polypeptides they

CC encode are useful for manipulating crop plants to alter or improve

CC phenotypic characteristics, to produce large quantities of oil or

CC proteins, to incur resistance to insecticides, viruses or fungi, and to

CC incur stress tolerance (e.g. salt, cold or drought) to ensure the plants

CC have a high nutritional value with reduced apical dominance or dwarfism,

CC early flowering or altered metabolic pathways. This sequence represents a

CC plant nucleic acid of the invention. Note: The sequence data for this

CC patent did not form part of the printed specification but was obtained in

CC electronic format directly from USPTO at seqdata.uspto.gov/sequence.html.

XX

XX Sequence 922 BP; 210 A; 294 C; 234 G; 184 T; 0 U; 0 Other;

QQ

Query Match 79.2%; Score 19.8; DB 12; Length 922;

Best Local Similarity 91.3%; Pred. No. 64;

Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24

Db 869 GAAGAAATTCAGTTCATAGCTTG 847

RESULT 2

AAA75080/C

ID AAA75080 standard; DNA; 44848 BP.

XX

AC AAA75080;

XX

DT 15-JAN-2001 (first entry)

XX

DE Nucleotide sequence of the human heparanase gene.

XX

XX Human; heparanase; gene therapy; tumour; inflammation; autoimmunity;

KW heparin-binding growth factor; cytokine; neurodegenerative plaque;

KW wound healing; infection; burn; angiogenesis; restenosis;

KW atherosclerosis; inflammation; neurodegenerative disease;

KW Gerstmann-Straussler Syndrome; Creutzfeldt-Jakob disease; ds.

XX

OS Homo sapiens.

XX

XX WO200052178-A1.

XX

XX 08-SEP-2000.

XX

PF 14-FEB-2000; 2000WO-US003542.

XX

PR 01-MAR-1999; 99US-00258892.

XX

XX (INSI-) INSIGHT STRATEGY & MARKETING LTD.

PA (HADA-) HADASIT MEDICAL RES SERVICES & DEV.

PA (FRIE/) FRIEDMAN M M.

XX

XX Pecker I, Vlodavsky I, Feinstein E;

XX WPI; 2000-579289/54.

DR

XX New polynucleotides encoding a polypeptide having heparanase activity,

PT useful in wound healing and in gene therapy, particularly in treating

PT tumor, inflammation, autoimmunity, neurodegenerative diseases.

XX

XX Claim 9; Page 131-143; 152pp; English.

XX

XX The present sequence represents a human gene which encodes a protein with

CC heparanase catalytic activity. The heparanase (hpa) polynucleotide is

CC useful in gene therapy, particularly in treating tumour, inflammation or

CC autoimmunity. Particularly, the polynucleotide is useful in modulating

CC the bioavailability of heparin-binding growth factors, cellular responses

CC to heparin-binding growth factors (e.g. bFGF) and cytokines (e.g.

CC interleukin (IL)-8), cell interaction with plasma lipoproteins, cellular

CC susceptibility to certain viral and some bacterial and protozoa

CC infections, or disintegration of neurodegenerative plaques. The

CC polynucleotide is also useful in wound healing (e.g. thermal, chemical or

CC radiation burns), and in the treatment of angiogenesis, restenosis,

CC atherosclerosis, inflammation, neurodegenerative diseases (Gerstmann-

CC Strausler Syndrome or Creutzfeldt-Jakob disease), and some viral,

CC bacterial or protozoa infections

XX

XX Sequence 44848 BP; 12560 A; 9646 C; 8930 G; 13712 T; 0 U; 0 Other;

QQ

Query Match 79.2%; Score 19.8; DB 3; Length 44848;

Best Local Similarity 91.3%; Pred. No. 94;

Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24

Db 30803 GAAGAAATTCAGTTCATAGCTTG 30781

RESULT 3

ADG88832/C

ID ADG88832 standard; DNA; 44848 BP.

XX

AC ADG88832;

XX

DT 11-MAR-2004 (first entry)

XX

DE Human hpa genomic DNA.

XX

KW Wound healing; heparanase; ulcer; burn; laceration; surgical incision;

KW necrosis; pressure wound; diabetic ulcer; angiogenesis; human; therapy;

KW gene; ss.

XX

OS Homo sapiens.

XX

XX Key Location/Qualifiers

FT CDS 2743..41863

FT /*tag= a

FT /product= "Human hpa protein"

XX

PN US2003161823-A1.

XX

PD 28-AUG-2003.

XX

XX 14-JAN-2003; 2003US-00341582.

XX

PR 31-AUG-1998; 98WO-US017954.

PR 01-MAR-1999; 99US-00258892.

PR 06-FEB-2001; 2001US-00776874.

PR 05-SEP-2001; 2001WO-IL000830.

PR 19-NOV-2001; 2001US-00988113.

XX

XX (ILAN/) ILAN N.

PA (VLOD/) VLODAVSKY I.

PA (YACO/) YACOBY-ZEEVI O.

PA (PECK/) PECKER I.

XX (FEIN/) FEINSTEIN E.

XX

XX Ilan N, Vlodavsky I, Yacoby-Zeevi O, Pecker I, Feinstein E;

```
XX WPI; 2003-897910/82.
DR P-PSDB; ADG88800.
XX
PT Composition for treating a wound comprising recombinant heparanase is
PT useful to induce or accelerate wound healing and induce or accelerate
PT angiogenesis.
XX
XX Claim 4; SEQ ID NO 42; 143pp; English.
XX
CC The present invention relates to methods and compositions for inducing
CC and/or accelerating wound healing via the catalytic activity of
CC heparanase. The invention is used to induce or accelerate a healing
CC process, particularly of an ulcer, burn, laceration, surgical incision,
CC necrosis, pressure wound, diabetic ulcer and to induce or accelerate
CC angiogenesis. The present sequence is human hpa genomic DNA.
XX
SQ Sequence 44848 BP; 12560 A; 9646 C; 8930 G; 13712 T; 0 U; 0 Other;
Query Match 79.2%; Score 19.8; DB 10; Length 44848;
Best Local Similarity 91.3%; Pred. No. 94;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 2 GAAGAAATTCAGTTCATAGCTTG 24
Db 30803 GAAGAAATTCAGTTCATAGCTTG 30781
RESULT 4
ADL16411/c
ID ADL16411 standard; DNA; 44848 BP.
XX
AC ADL16411;
XX
XX 06-MAY-2004 (first entry)
XX Human heparanase genomic DNA.
DE
XX Human; ds; heparanase; gene; heparanase-dependent cancer; cancer;
XX autoimmune reaction; inflammation; chromosome 4.
XX Homo sapiens.
XX
FH Key Location/Qualifiers
FT promoter 2535..2635
FT /*tag= a
FT /note= "Minimal promoter region. Claimed in claim 15"
FT 5'UTR 2635..2742
FT /*tag= b
FT /note= "Claimed in claim 19"
FT 3'UTR 41864..41890
FT /*tag= c
FT /note= "Claimed in claim 20"
XX
XX US2003236215-A1.
XX
XX 25-DEC-2003.
XX
XX 09-JUN-2003; 2003US-00456573.
XX
XX 31-AUG-1998; 98WO-US017954.
XX 01-MAR-1999; 99US-00258892.
XX 08-NOV-1999; 99US-00435739.
XX
XX (INSIGHT STRATEGY & MARKETING LTD.
PA (HADA-) HADASIT MEDICAL RES SERVICES & DEV.
XX
XX Pecker I, Vlodavsky I, Feinstein B;
XX
XX WPI; 2004-070610/07.
XX
XX New antisense oligonucleotide hybridizable with a polynucleotide encoding
PT a polypeptide with heparanase activity, useful for treating diseases such
```

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as cancer and autoimmune disorders.
Claim 2; SEQ ID NO 42; 108pp; English.
XX
XX The invention relates to an antisense oligonucleotide (ASO) comprising a
XX polynucleotide or a polynucleotide analogue of at least 10 bases being
XX hybridizable in vivo, under physiological conditions, with a portion of
XX a polynucleotide strand encoding a polypeptide having heparanase
XX catalytic activity. Also included are a method of in vivo downregulating
XX heparanase activity (comprising administering the ASO in vivo), a method
XX of treating a subject suffering from a pathological condition
XX (characterised by heparanase activity, comprising administering ASO to
XX the subject), a pharmaceutical composition comprising the ASO and a
XX carrier, an antisense nucleic acid construct (comprising a promoter
XX sequence and a polynucleotide sequence directing the synthesis of an
XX antisense RNA sequence of at least 10 bases being hybridizable in vivo,
XX under physiological conditions, with a polynucleotide strand encoding a
XX polypeptide having heparanase catalytic activity), a method of in vivo
XX downregulating heparanase activity (comprising administering in vivo the
XX antisense nucleic acid construct), a pharmaceutical composition
XX comprising the antisense nucleic acid construct and a carrier, and an
XX antisense oligonucleotide comprising a polynucleotide or a polynucleotide
XX analogue of at least 10 bases being hybridizable in vivo, under
XX physiological conditions, with a portion of a polynucleotide strand being
XX characterised by forming at least a portion of an untranslated region
XX (UTR) for a polynucleotide strand encoding a polypeptide having
XX heparanase catalytic activity. The methods and compositions of the
XX present invention are useful for the prevention and/or treatment of
XX diseases or conditions associated with aberrant heparanase activity, such
XX as heparanase-dependent cancer, cancer, autoimmune reaction and
XX inflammation. The gene for human heparanase is located on chromosome 4.
XX The present sequence is the human heparanase gene.
XX
SQ Sequence 44848 BP; 12560 A; 9646 C; 8930 G; 13712 T; 0 U; 0 Other;
Query Match 79.2%; Score 19.8; DB 12; Length 44848;
Best Local Similarity 91.3%; Pred. No. 94;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 2 GAAGAAATTCAGTTCATAGCTTG 24
Db 30803 GAAGAAATTCAGTTCATAGCTTG 30781
RESULT 5
ADM48748/c
ID ADM48748 standard; DNA; 44848 BP.
XX
AC ADM48748;
XX
XX 03-JUN-2004 (first entry)
XX Human hpa genomic DNA.
XX
XX Transgenic animal; heparanase; cancer; viral infection; restenosis;
XX neurodegenerative disease; atherosclerosis; pulmonary disorder; hpa;
XX human; gene; ds.
XX Homo sapiens.
XX
XX Key Location/Qualifiers
XX exon 2635..2969
XX /*tag= a
XX /number= 1
XX CDS 2743..41863
XX /*tag= b
XX /product= "Hpa protein"
XX intron 2970..15141
XX /*tag= c
XX /number= 1
XX exon 15142..15287
XX /*tag= d
XX /number= 2
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FT intron 18162..24214
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FT intron 26770..27413
FT      /*tag= k
FT      /number= 5
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FT      /*tag= l
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FT exon 31174..31288
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FT      /number= 9
FT intron 31289..35227
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FT      /number= 9
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FT intron 35347..36389
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FT      /number= 10
FT exon 36390..36536
FT      /*tag= v
FT      /number= 11
FT intron 36537..41703
FT      /*tag= w
FT      /number= 11
FT exon 41704..41890
FT      /*tag= x
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FT      /number= 12
XX US2003217375-A1.
XX
XX 20-NOV-2003.
XX
XX 24-FEB-2003; 2003US-00371218.
XX
XX 31-AUG-1998; 98WO-US017954.
XX 01-MAR-1999; 99US-00258892.
XX 06-FEB-2001; 2001US-00776874.
XX 19-NOV-2001; 2001US-00988113.
XX
XX (ZCHA/) ZCHARIA E.
PA
PA (VL0D/) VLODAVSKY I.
PA (METZ/) METZGER S.
PA (PECK/) PECKER I.
PA (ILAN/) ILAN N.
PA (CHAJ/) CHAJEK-SHAUL T.
PA (GOLD/) GOLDSHMIDT O.
XX
XX Zcharia E, Vlodayvsky I, Metzger S, Pecker I, ilan N;
XX Chajek-Shaul T, Goldshmidt O;
XX WPI; 2004-021918/02.
XX P-PSDB; ADM48759.
XX
XX New transgenic non-human animal expressing heparinase, useful as models
XX for human disease, such as cancers, viral infection, neurodegenerative
XX diseases, restenosis, atherosclerosis and pulmonary disorders.
XX
XX Example 10; SEQ ID NO 42; 106pp; English.
XX
XX The present invention relates to a transgenic non-human animal whose
XX genome comprises an exogenous polynucleotide sequence, including a
XX promoter active in tissues of the non-human, a region encoding a human
XX heparanase, where the promoter and the region encoding human heparanase
XX are operably linked in the exogenous polynucleotide such that human
XX heparanase is expressed in at least a portion of the cells of the non-
XX human animal. The methods and compositions of the present invention are
XX useful for the production of transgenic animals expressing heparanase, to
XX be used as models for human diseases such as cancers, viral infection,
XX restenosis, neurodegenerative diseases, atherosclerosis and pulmonary
XX disorders. The present sequence is human hpa genomic DNA used in the
XX exemplification of the invention.
XX
XX Sequence 44848 BP; 12560 A; 9646 C; 8930 G; 13712 T; 0 U; 0 Other;
SQ
Query Match 79.2%; Score 19.8; DB 12; Length 44848;
Best Local Similarity 91.3%; Pred. No. 94;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 2 GAAGAAATTCAGTTCATAGCTTG 24
Db 30803 GAAGAAATTCAGTTCATAGCTTG 30781
RESULT 6
ADL13873/c
ID ADL13873 standard; DNA; 183178 BP.
XX
XX AC ADL13873;
XX
XX DT 06-MAY-2004 (first entry)
XX
XX DE Osteoarthritis-associated polymorphic nucleotide #405.
XX
XX KW ds; gene; osteopathic; antiinflammatory; antiarthritic; gene therapy;
XX joint space narrowing; osteophyte development; joint pain;
XX osteoarthritis; SNP; single nucleotide polymorphism.
XX
XX OS Homo sapiens.
XX
XX PN WO2003054166-A2.
XX
XX PD 03-JUL-2003.
XX
XX PF 19-DEC-2002; 2002WO-US041225.
XX
XX PR 20-DEC-2001; 2001US-0342603P.
XX
XX PA (INCY-) INCYTE GENOMICS INC.
XX
XX PI Jones KA, Schafer A;
XX
XX DR WPI; 2003-559141/52.
XX
```

PT Determining susceptibility of an individual to joint space narrowing,
 PT osteophyte development and/or joint pain comprises identifying whether
 PT the individual has at least one polymorphism in a polynucleotide encoding
 PT a protein.

PS Disclosure; SEQ ID NO 405; 297pp; English.

XX The invention relates to a method of determining susceptibility of an
 CC individual to joint space narrowing and/or osteophyte development and/or
 CC joint pain comprising identifying whether the individual has at least one
 CC polymorphism in a polynucleotide encoding at least one of the protein
 CC listed in the specification. The methods, composition and agent are
 CC useful for modulating the susceptibility of an individual to joint space
 CC narrowing and/or osteophyte development and/or joint pain that is
 CC associated with a disease, preferably osteoarthritis. The cell line and
 CC the non-human animal are useful for screening for an agent for diagnosing
 CC an individual having susceptibility to joint space narrowing and/or
 CC osteophyte development and/or joint pain. This sequence corresponds to
 CC the polynucleotide encoding a protein listed in the specification. (Note:
 CC The sequence data for this patent did not form part of the printed
 CC specification but was obtained in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences).

SQ Sequence 183178 BP; 52771 A; 36295 C; 36571 G; 54082 T; 0 U; 3459 Other;

Query Match 79.2%; Score 19.8; DB 10; Length 183178;

Best Local Similarity 91.3%; Pred. No. 1.1e+02; Indels 0; Gaps 0;
 Matches 21; Conservative 0; Mismatches 2;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24

DB 155952 GAAGAAATTCAGTTCATAGCTTG 155930

RESULT 7

ID ACN44572 standard; DNA; 49507 BP.

AC ACN44572;

DT 18-NOV-2004 (first entry)

XX Mouse genomic sequence MCG21659.

XX Cytostatic; carcinoma; lymphoma; cancer; murine; gene; ss.

XX Mus musculus.

XX WO2003073826-A2.

XX 12-SEP-2003.

XX 28-FEB-2003; 2003WO-US0006235.

XX 01-MAR-2002; 2002US-00087192.

XX (SAGR-) SAGRES DISCOVERY.

XX Morris DW;

XX WPI; 2003-328604/31.

XX Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
 PT comprises a nucleotide sequence.

PS Claim 1; SEQ ID NO 1087; 0pp; English.

XX The present invention relates to novel DNA and protein sequences which
 CC are associated with carcinomas. The sequences are useful for: (i) for
 CC screening drug candidates; (ii) for screening of bioactive agent capable
 CC of binding to carcinoma Associated protein (CAP); (iii) for screening of
 CC a bioactive agent capable of modulating the activity of CAP; (iv) for
 CC evaluating the effect of a candidate carcinoma drug; (v) for diagnosing

CC carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
 CC carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
 CC (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
 CC determining Carcinoma Associated (CA) gene copy number. In addition, the
 CC CA genes are useful as DNA vaccines and the CAP are useful as markers of
 CC carcinoma including lymphoma. The present sequence is one such CA coding
 CC sequence. Note: this patent is an equivalent to basic patent
 CC US2002182586A1, for which no sequence data was published

XX SQ Sequence 49507 BP; 14131 A; 9178 C; 9693 G; 15785 T; 0 U; 720 Other;

Query Match 76.8%; Score 19.2; DB 11; Length 49507;

Best Local Similarity 87.5%; Pred. No. 1.8e+02; Indels 0; Gaps 0;
 Matches 21; Conservative 0; Mismatches 3;

QY 1 TGAAGAAATTCAGTTCATAGCTTG 24

DB 1473 TGAAGAAATTCAGTTCATAGCTTG 1496

RESULT 8

ABS52506/c

ID ABS52506 standard; DNA; 202001 BP.

XX ABS52506;

XX 15-NOV-2002 (first entry)

XX Human transporter protein genomic DNA.

XX Human; gene; ds; transporter protein; cell proliferation;

KW cell differentiation; cell signaling; sodium bicarbonate cotransporter;
 KW transgenic; ligand transport; drug development; SNP;
 KW single nucleotide polymorphism.

XX Homo sapiens.

XX Key Location/Qualifiers

FT variation replace(2180,C)
 FT /tag= ax
 FT /standard_name= "Single nucleotide polymorphism"

FT exon 3016..3096

FT /tag= a

FT intron 3097..11617

FT /tag= b

FT /number= 1

FT /cons_splice= (5'site:yes,3'site:no)

FT variation replace(4693,C)

FT /tag= ay

FT exon 11618..11690

FT /tag= c

FT intron 11691..37943

FT /tag= d

FT /number= 2

FT /cons_splice= (5'site:no,3'site:no)

FT variation replace(13759,T)

FT /tag= az

FT variation replace(17580,A)

FT /tag= ba

FT variation replace(17701,A)

FT /tag= bc

FT variation replace(18151,C)

FT /tag= bd

FT variation replace(21076,C)

FT /tag= be

FT /standard_name= "Single nucleotide polymorphism"

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FT replace(25961, .25962,TWT)
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FT replace(29242,T)
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FT /standard_name= "Single nucleotide polymorphism"
FT replace(35888,A)
FT /*tag= bm
FT /standard_name= "Single nucleotide polymorphism"
FT replace(37779,A)
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FT /number= 3
FT 38091, .76221
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FT replace(72117,G)
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FT /standard_name= "Single nucleotide polymorphism"
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FT /number= 4
FT 76361, .91402
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FT /cons_splice= (5'site:yes,3'site:no)
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FT replace(77426,G)
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FT replace(89179,G)
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FT /standard_name= "Single nucleotide polymorphism"
FT replace(90081,C)
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FT 91403, .91563
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FT replace(96808,C)
FT /*tag= ci
FT /standard_name= "Single nucleotide polymorphism"
FT 99312, .99500
FT /*tag= k
FT /number= 6
FT 99501, .108867
FT /*tag= l
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FT /standard_name= "Single nucleotide polymorphism"
FT replace(105911,T)
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FT /standard_name= "Single nucleotide polymorphism"
FT 108868, .108959
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FT 108960, .110489
FT /*tag= n
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FT      Best Local Similarity 87.58; Pred.No.2e+02;
FT      Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY      1 TGAAGAAATTCAGTTCATAGCTTG 24
Db      174078 TGAAGAAATTCATTCAAGCTTG 174055
RESULT 9
ADG46742/c
ID      ADG46742 standard; DNA; 202001 BP.
XX
AC      ADG46742;
XX
DT      11-MAR-2004 (first entry)
XX
DE      Human transporter genomic DNA.
XX
KW      Transporter protein; therapy; human; chromosome 2; gene; ds.
XX
OS      Homo sapiens.
XX
FH      Key
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FT      3016..3096
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FT      3097..11617
FT      /*tag= d
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FT      11691..37943
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FT      replace(65909,A)
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FT replace(114256,A)
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FT 115864..118931
FT /*tag= bp
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Query Match 76.8%; Score 19.2; DB 10; Length 202001;
Best Local Similarity 87.5%; Pred. No. 2e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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QY 1 TGAAGAAATTCAGTTCATAGCTTG 24
DB 174078 TGAAGAAATTCATTTCAAAGCTTG 174055
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RESULT 10

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ADP92027
ID ADP92027 standard; cDNA; 235 BP.
XX
AC ADP92027;
XX
DT 09-SEP-2004 (first entry)
XX
DE Cotton expressed sequence tag, EST, #1038.
XX
KW Cotton; ss; EST; expressed sequence tag; plant; plant protection;
KW plant improvement; marker-assisted breeding.
XX
OS Gossypium hirsutum; variety Nucleotid33B.
XX
PN US2004123338-A1.
XX
PD 24-JUN-2004.
XX
PF 08-DEC-2000; 2000US-00732627.
XX
PR 10-DEC-1999; 99US-0170255P.
XX (FINC/) FINCHER K L.
XX PI Fincher KL;
XX WPI; 2004-479807/45.
XX
PT New substantially purified nucleic acid molecule that encodes a cotton
PT protein or its fragment, useful as molecular tool for the targeting and
PT isolation of novel genes for plant protection and improvement.
XX
PS Claim 1; SEQ ID NO 1038; 30pp; English.
XX
CC The invention relates to a substantially purified nucleic acid molecule
CC that encodes a cotton protein or its fragment comprising an EST
CC (expressed sequence tag) appearing as ADP9090-ADP95919. Also included
CC are a substantially purified cotton protein or its fragment encoded by a
CC nucleic acid molecule above and a transformed plant (having a nucleic
CC acid molecule which comprises: an exogenous promoter region which
CC functions in a plant cell to cause the production of a mRNA molecule; a
CC structural nucleic acid molecule comprising one of the ESTs or their
CC complements; a 3' non-translated sequence that functions in the plant
CC cell to cause termination of transcription and addition of polyadenylated
CC ribonucleotides to a 3' end of the mRNA molecule). The ESTs are useful as
CC molecular tool for the targeting and isolation of novel genes for plant
CC protection and improvement. The ESTs are useful for developing new
CC strategies for understanding critical plant developmental and metabolic
CC pathways, for isolating genes and promoters, for identifying and mapping
CC the genes involved in developmental and metabolic pathways, and for
CC determining gene function. The cotton nucleic acid molecules are useful
CC as molecular tags to isolate genetic regions, isolate genes, map genes,
CC and determine gene function. The nucleic acid molecules are useful for
CC determining if genes are members of a particular gene family and for use
CC in marker-assisted breeding programs. The present sequence is one of the
CC 4930 cotton ESTs of the invention. NOTE: The sequences are not displayed
CC in the specification but are available in electronic format from the
CC USPTO at seqdata.uspto.gov/sequence.html?DocID=20040123338.
XX
SQ Sequence 235 BP; 66 A; 40 C; 60 G; 69 T; 0 U; 0 Other;
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Query Match 76.0%; Score 19; DB 12; Length 235;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 TGAAGAAATTCAGTTCATA 19
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Db      179 TGAAGAAATTCAGTTTCATA 197
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RESULT 11
ACH97547/c
ID ACH97547 standard; DNA; 1227 BP.
XX AC ACH97547;
XX DT 29-JUL-2004 (first entry)
XX DE Klebsiella pneumoniae polynucleotide seqid 3342.
XX KW Recombinant expression vector; transcription regulatory element;
XX KW Klebsiella pneumoniae protein; antibacterial; Vaccine; gene; ds.
XX OS Klebsiella pneumoniae.
XX PN US6610836-B1.
XX PD 26-AUG-2003.
XX PF 27-JAN-2000; 2000US-00489039.
XX PR 29-JAN-1999; 99US-0117747P.
XX PA (GENO-) GENOME THERAPEUTICS CORP.
XX PI Breton GL, Osborne M;
XX DR WPI; 2003-895346/82.
XX DR P-PSDB; ABO63996.
XX PT New nucleic acid encoding a Klebsiella pneumoniae polypeptide, useful for
XX PT preparing a vaccine composition against Klebsiella pneumoniae.
XX PS Disclosure; SEQ ID NO 3342; 932pp; English.
XX CC The invention describes a new isolated nucleic acid encoding a Klebsiella
XX CC pneumoniae polypeptide. Also described are: a recombinant expression
XX CC vector comprising the nucleic acid, operably linked to a transcription
XX CC regulatory element; and a cell comprising the recombinant expression
XX CC vector. The nucleic acid is useful for preparing a vaccine composition
XX CC against Klebsiella pneumoniae. This sequence encodes a Klebsiella
XX CC pneumoniae polypeptide of the invention
XX SQ Sequence 1227 BP; 222 A; 351 C; 360 G; 294 T; 0 U; 0 Other;

Query Match 75.2%; Score 18.8; DB 11; Length 1227;
Best Local Similarity 90.9%; Pred. No. 1.8e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTTCATAGCT 22
|||||
Db 937 TGATGAATCCAGTTTCATAGCT 916

RESULT 12
ADQ97167/c
ID ADQ97167 standard; DNA; 176771 BP.
XX AC ADQ97167;
XX DT 07-OCT-2004 (first entry)
XX DE Human cancer associated sequence HD2-08-009, SEQ ID 143.
XX KW Cytostatic; Gene Therapy; cancer; leukemia; lymphoma; Human; ds.
XX OS Homo sapiens.
XX PS WO2004060304-A2.

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XX 22-JUL-2004.
XX 22-DEC-2003; 2003WO-US041389.
XX 27-DEC-2002; 2002US-00330773.
XX (SAGR-) SAGRES DISCOVERY INC.
XX Morris DW, Malandro MS;
XX WPI; 2004-543781/52.
XX New isolated cancer associated nucleic acids comprising at least 10
XX PT contiguous nucleotides, useful for diagnosing, preventing and/or treating
XX PT cancers such as leukemia and lymphoma.
XX PS Claim 1; SEQ ID NO 143; 199pp; English.
XX CC The present invention relates to cancer associated sequences (ADQ97025-
XX CC ADQ98004). The sequences are useful for the diagnosis, prevention and/or
XX CC treatment of cancer, such as leukemia and lymphoma. Note: The sequence
XX CC data for this patent did not form part of the printed specification, but
XX CC was obtained in electronic format directly from WIPO at
XX CC ftp.wipo.int/pub/published_pct_sequences.
XX SQ Sequence 176771 BP; 47665 A; 34089 C; 38342 G; 56655 T; 0 U; 20 Other;

Query Match 75.2%; Score 18.8; DB 12; Length 176771;
Best Local Similarity 90.9%; Pred. No. 3e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTTCATAGCT 22
|||||
Db 100466 TGAGAAATTCATTATAGCT 100445

RESULT 13
ABZ14865
ID ABZ14865 standard; DNA; 4212 BP.
XX AC ABZ14865;
XX DT 21-JAN-2003 (first entry)
XX DE Arabidopsis thaliana stress regulated gene SEQ ID NO 2670.
XX KW Arabidopsis thaliana; plant; gene; stress; transgenic; ds.
XX OS Arabidopsis thaliana.
XX PN WO200216655-A2.
XX PD 28-FEB-2002.
XX PF 24-AUG-2001; 2001WO-US026685.
XX PR 24-AUG-2000; 2000US-0227866P.
XX PR 26-JAN-2001; 2001US-0264647P.
XX PR 22-JUN-2001; 2001US-0300111P.
XX PA (SCRI ) SCRIPPS RES INST.
XX PA (SYGN ) SYNGENTA PARTICIPATIONS AG.
XX PI Harper JF, Kreps J, Wang X, Zhu T;
XX WPI; 2002-304127/34.
XX PT Identifying a stress condition to which a plant cell has been exposed and
XX PT producing plants with increased tolerance to these abiotic stresses.
XX PS Claim 144; SEQ ID NO 2670; 577pp + Sequence Listing; English.

```

CC The invention relates to identifying a stress condition to which a plant
 CC cell has been exposed, comprising: (a) contacting nucleic acid with an
 CC representative of expressed polynucleotides in the plant cell with an
 CC array or probes representative of the plant cell genome; and (b)
 CC detecting a profile of expressed polynucleotides in the plant cell
 CC characteristic of a stress response. The method is useful in the
 CC production of transgenic plants, cells and seeds and in producing plants
 CC with increased tolerance to abiotic stress. The present sequence is that
 CC of an Arabidopsis thaliana stress regulated gene (AB212196-AB217574) used
 CC in methods of the invention. Note: The sequence data for this patent is
 CC not represented in the printed specification but is based on sequence
 CC information supplied to Derwent by the European Patent Office
 XX

SQ Sequence 4212 BP; 1235 A; 892 C; 963 G; 1122 T; 0 U; 0 Other;

Query Match 74.4%; Score 18.6; DB 6; Length 4212;
 Best Local Similarity 84.0%; Pred. No. 2.6e+02;
 Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTCATAGCTTGT 25
 Db 2453 TGAAGAAAGTTAGCTCATCTTGT 2477

RESULT 14
 ACN44212
 ID ACN44212 standard; DNA; 116858 BP.

XX ACN44212;

DT 18-NOV-2004 (first entry)

DE Mouse genomic sequence MCG15964.

KW Cytostatic; carcinoma; lymphoma; cancer; murine; gene; ss.

OS Mus musculus.

PN WO2003073826-A2.

PD 12-SEP-2003.

PF 28-FEB-2003; 2003WO-US006235.

PR 01-MAR-2002; 2002US-00087192.

XX (SAGR-) SAGRES DISCOVERY.

PA Morris DW;

DR WPI; 2003-328604/31.

XX Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
 PT comprises a nucleotide sequence.

PS Claim 1; SEQ ID NO 547; Opp; English.

XX The present invention relates to novel DNA and protein sequences which
 CC are associated with carcinomas. The sequences are useful for: (i) for
 CC screening drug candidates; (ii) for screening of bioactive agent capable
 CC of binding to Carcino Associated Protein (CAP); (iii) for screening of
 CC a bioactive agent capable of modulating the activity of CAP; (iv) for
 CC evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
 CC carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
 CC carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
 CC (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
 CC determining Carcino Associated (CA) gene copy number. In addition, the
 CC CA genes are useful as DNA vaccines and the CAP are useful as markers of
 CC carcinoma including lymphoma. The present sequence is one such CA coding
 CC sequence. Note: This patent is an equivalent to basic patent
 CC US2002182586A1, for which no sequence data was published
 XX

SQ Sequence 116858 BP; 32402 A; 22234 C; 23982 G; 34772 T; 0 U; 3468 Other;

Query Match 74.4%; Score 18.6; DB 11; Length 116858;
 Best Local Similarity 84.0%; Pred. No. 3.5e+02;
 Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTCATAGCTTGT 25
 Db 17178 TGAAGAAATTCATATATTTTGT 17202

RESULT 15
 ADK54409/c

ID ADK54409 standard; DNA; 978 BP.

XX ADK54409;

DT 06-MAY-2004 (first entry)

DE Plant DNA sequence which confers altered metabolic characteristic #1792.

KW altered metabolic characteristic; plant; acid metabolism;

KW alcohol metabolism; fatty acid metabolism;

KW branched fatty acid metabolism; alkaloid metabolism;

KW amino acid metabolism; ester metabolism; glyceride metabolism;

KW phenolic metabolism; carbohydrate metabolism; sterol metabolism;

KW terpene metabolism; isoprenoid metabolism; alkene metabolism;

KW alkyne metabolism; hydrocarbon metabolism; ketone metabolism;

KW quinone metabolism; disease resistance; gene shuffling; sexual PCR; ds.

XX Unidentified.

XX WO2003020936-A1.

XX 13-MAR-2003.

XX 30-AUG-2002; 2002WO-US027894.

XX 31-AUG-2001; 2001US-0316471P.

XX (DOWC) DOW CHEM CO.

XX (DOWC) DOW AGROSCIENCES LLC.

PI Weglarz T, Gachotte D, Blakeslee B, McCreary DA, Pell RJ;

PI Orledo JVB, Crosley R, Reddy AS, Shukla V, Larrinua I, Miller BA;

XX WPI; 2003-313091/30.

XX Novel genes that confer altered metabolic characteristics in Nicotiana
 PT benthamiana plants, useful for altering the levels of metabolites e.g.
 PT acids, fatty acids, amino acids, carbohydrates, hydrocarbons and sterols.

PS Claim 1; SEQ ID NO 1792; 2576pp; English.

XX The invention comprises DNA sequences which confer an altered metabolic
 CC characteristic when they are expressed in a plant. The DNA sequences of
 CC the invention are useful for producing plants with an altered metabolic
 CC characteristic, such as: altered acid metabolism, alcohol metabolism,
 CC fatty acid metabolism, branched fatty acid metabolism, alkaloid or other
 CC base metabolism, altered amino acid metabolism, altered ester metabolism,
 CC altered glyceride metabolism, altered phenolic metabolism, altered
 CC carbohydrate metabolism, altered sterol, oxygenated terpene, or
 CC isoprenoid metabolism, alkene or alkyne metabolism, hydrocarbon
 CC metabolism, ketone or quinone metabolism. The DNA sequences of the
 CC invention may be used to provide disease resistance in a plant and gene
 CC shuffling or sexual PCR procedures. The present nucleic acid represents a
 CC DNA sequence of the invention.

SQ Sequence 978 BP; 358 A; 146 C; 164 G; 310 T; 0 U; 0 Other;

Query Match 72.8%; Score 18.2; DB 10; Length 978;
 Best Local Similarity 87.0%; Pred. No. 3.3e+02;
 Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 AAGAAATTCAGTTCATAGCTTGT 25
|||
Db 801 AATAAATTCAGTTCAGTTCATAGCTTGT 779

Search completed: August 13, 2005, 04:14:41
Job time : 206.266 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: August 13, 2005, 03:33:32 ; Search time 61.4362 Seconds
(without alignments)
665.844 Million cell updates/sec

Title: US-10-673-854-3

Perfect score: 25

Sequence: 1 tgaagaattcagttcagttcgtgt 25

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA.*

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2: /cgn2_6/ptodata/1/ina/5B COMB.seq.*

3: /cgn2_6/ptodata/1/ina/6A COMB.seq.*

4: /cgn2_6/ptodata/1/ina/6B COMB.seq.*

5: /cgn2_6/ptodata/1/ina/PCTUS COMB.seq.*

6: /cgn2_6/ptodata/1/ina/backfileseq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	19.8	79.2	44848	US-09-435-739-42	Sequence 42, Appl
C 2	19.8	79.2	44848	US-09-988-113-42	Sequence 42, Appl
C 3	19.2	76.8	202001	US-09-734-674-3	Sequence 3, Appli
C 4	18.8	75.2	1227	US-09-489-039A-3342	Sequence 3342, Ap
C 5	18.2	72.8	29326	US-09-949-016-15356	Sequence 15356, A
C 6	18.2	72.8	29720	US-09-949-016-16521	Sequence 16521, A
7	18.2	72.8	126176	US-09-949-016-16137	Sequence 16137, A
8	18.2	72.8	126176	US-09-949-016-16138	Sequence 16138, A
C 9	18.2	72.8	223471	US-09-949-016-12387	Sequence 12387, A
C 10	18.2	72.8	223471	US-09-949-016-12724	Sequence 12724, A
C 11	18.2	72.8	223471	US-09-949-016-12725	Sequence 12725, A
12	18	72.0	447	US-09-621-976-17257	Sequence 17257, A
13	17.8	71.2	601	US-09-949-016-157433	Sequence 157433, A
14	17.8	71.2	601	US-09-949-016-157540	Sequence 157540, A
C 15	17.8	71.2	1325	US-09-710-279-999	Sequence 999, App
C 16	17.8	71.2	1485	US-09-134-001C-1339	Sequence 1339, Ap
17	17.8	71.2	2978	US-09-710-279-4039	Sequence 4039, Ap
18	17.8	71.2	3745	US-09-710-279-4045	Sequence 4045, Ap
19	17.6	70.4	41988	US-09-949-016-14501	Sequence 14501, A
20	17.6	70.4	77618	US-09-949-016-11768	Sequence 11768, A
21	17.6	70.4	119153	US-09-949-016-12378	Sequence 12378, A
22	17.4	69.6	601	US-09-949-016-179792	Sequence 179792, A
23	17.4	69.6	601	US-09-949-016-179793	Sequence 179793, A
24	17.4	69.6	390416	US-09-348-352-1292	Sequence 1292, Ap
25	17.2	68.8	192	US-09-328-352-1292	Sequence 1292, Ap
26	17.2	68.8	601	US-09-949-016-93041	Sequence 93041, A
27	17.2	68.8	601	US-09-949-016-93042	Sequence 93042, A

C 28	17.2	68.8	601	US-09-949-016-153121	Sequence 153121, A
C 29	17.2	68.8	601	US-09-949-016-153122	Sequence 153122, A
C 30	17.2	68.8	601	US-09-949-016-153123	Sequence 153123, A
C 31	17.2	68.8	601	US-09-949-016-177341	Sequence 177341, A
32	17.2	68.8	601	US-09-949-016-177342	Sequence 177342, A
33	17.2	68.8	601	US-09-949-016-177343	Sequence 177343, A
34	17.2	68.8	601	US-09-949-016-177344	Sequence 177344, A
35	17.2	68.8	2521	US-09-620-312D-1015	Sequence 1015, Ap
36	17.2	68.8	2250	US-09-620-312D-1014	Sequence 16808, A
37	17.2	68.8	13438	US-09-949-016-16808	Sequence 16808, A
38	17.2	68.8	35058	US-09-949-016-12607	Sequence 12607, A
39	17.2	68.8	35059	US-09-949-016-13831	Sequence 13831, A
C 40	17.2	68.8	74177	US-09-949-016-11988	Sequence 11988, A
C 41	17.2	68.8	74177	US-09-949-016-17388	Sequence 17388, A
C 42	17.2	68.8	83697	US-09-949-016-16040	Sequence 16040, A
C 43	17.2	68.8	126468	US-09-949-016-14418	Sequence 14418, A
44	17.2	68.8	131724	US-09-949-016-12893	Sequence 12893, A
C 45	17.2	68.8	176006	US-09-949-016-16804	Sequence 16804, A

ALIGNMENTS

RESULT 1

US-09-435-739-42/c

; Sequence 42, Application US/09435739

; Patent No. 6664105

; GENERAL INFORMATION:

; APPLICANT: Pecker, Iris

; APPLICANT: Vlodavsky, Israel

; APPLICANT: Feinstein, Elena

; TITLE OF INVENTION: POLYNUCLEOTIDE ENCODING A POLYPEPTIDE HAVING HEPARANASE ACTIVITY

; TITLE OF INVENTION: EXPRESSION OF SAME IN GENETICALLY MODIFIED CELLS

; FILE REFERENCE: 00/20454

; CURRENT APPLICATION NUMBER: US/09/435,739

; CURRENT FILING DATE: 2001-06-05

; NUMBER OF SEQ ID NOS: 47

; SOFTWARE: PatentIn version 3.0

; SEQ ID NO 42

; LENGTH: 44848

; TYPE: DNA

; ORGANISM: Homo sapiens

; US-09-435-739-42

Query Match 79.2%; Score 19.8; DB 4; Length 44848;

Best Local Similarity 91.3%; Pred. No. 32;

Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GAAGAAATTCAGTTTCATAGCTTG 24

Db 30803 GAAGAAATTCAGTTTCATAGCTTG 30781

RESULT 2

US-09-988-113-42/c

; Sequence 42, Application US/09988113

; Patent No. 6790658

; GENERAL INFORMATION:

; APPLICANT: Pecker, Iris

; APPLICANT: Vlodavsky, Israel

; APPLICANT: Feinstein, Elena

; TITLE OF INVENTION: POLYNUCLEOTIDE ENCODING A POLYPEPTIDE HAVING HEPARANASE ACTIVITY

; TITLE OF INVENTION: EXPRESSION OF SAME IN GENETICALLY MODIFIED CELLS

; FILE REFERENCE: 01/22781

; CURRENT APPLICATION NUMBER: US/09/988,113

; CURRENT FILING DATE: 2001-11-19

; PRIOR APPLICATION NUMBER: US 09/776,874

; PRIOR FILING DATE: 2001-02-06

; PRIOR APPLICATION NUMBER: US09/258,892

; PRIOR FILING DATE: 1999-03-01

; PRIOR APPLICATION NUMBER: PCT/US98/17954

; PRIOR FILING DATE: 1998-08-31

; PRIOR APPLICATION NUMBER: US 09/109,386

; PRIOR FILING DATE: 1998-07-02
; PRIOR APPLICATION NUMBER: US 08/922,170
; PRIOR FILING DATE: 1997-09-02
; NUMBER OF SEQ ID NOS: 47
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 42
; LENGTH: 44848
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-988-113-42

Query Match 79.2%; Score 19.8; DB 4; Length 44848;
Best Local Similarity 91.3%; Pred. No. 32;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
Db 30803 GAAGAAATTCAGTTCATAGCTTG 30781
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RESULT 3

US-09-734-674-3/c
; Sequence 3, Application US/09734674
; Patent No. 6498022
; GENERAL INFORMATION:
; APPLICANT: WEI, Ming-Hui et al
; TITLE OF INVENTION: ISOLATED HUMAN TRANSPORTER PROTEINS,
; TITLE OF INVENTION: NUCLEIC ACID MOLECULES ENCODING HUMAN TRANSPORTER PROTEINS,
; TITLE OF INVENTION: AND USES THEREOF
; FILE REFERENCE: CL001018
; CURRENT APPLICATION NUMBER: US/09/734,674
; CURRENT FILING DATE: 2000-12-13
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)...(202001)
; OTHER INFORMATION: n = A,T,C or G
US-09-734-674-3

Query Match 76.8%; Score 19.2; DB 4; Length 202001;
Best Local Similarity 87.5%; Pred. No. 70;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTCATAGCTTG 24
Db 174078 TGAAGAAATTCATTCAGGCTTG 174055
|||||

RESULT 4

US-09-489-039A-3342/c
; Sequence 3342, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Berton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; TITLE OF INVENTION: PNEUMONIAE FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A
; CURRENT FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 3342
; LENGTH: 1227
; TYPE: DNA
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-3342

Query Match 75.2%; Score 18.8; DB 4; Length 1227;
Best Local Similarity 90.9%; Pred. No. 58;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TGAAGAAATTCAGTTCATAGCT 22
Db 937 TGATGAATCCAGTTCATAGCT 916
|||||

RESULT 5

US-09-949-016-15356/c
; Sequence 15356, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15356
; LENGTH: 29326
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-15356

Query Match 72.8%; Score 18.2; DB 4; Length 29326;
Best Local Similarity 87.0%; Pred. No. 1.6e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
Db 11592 GAAGAAATTCAGTTCGCTTG 11570
|||||

RESULT 6

US-09-949-016-16521/c
; Sequence 16521, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16521
; LENGTH: 29720
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-16521

Query Match 72.8%; Score 18.2; DB 4; Length 29720;
Best Local Similarity 87.0%; Pred. No. 1.6e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24

Db 4957 GAAGGAATTCAGTTCATGCTTG 4935
|||||

RESULT 7
US-09-949-016-16137
; Sequence 16137, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16137
; LENGTH: 126176
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-16137

Query Match 72.8%; Score 18.2; DB 4; Length 126176;
Best Local Similarity 87.0%; Pred. No. 1.8e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||

Db 44879 GAAGAAATTCAGTTCCTAGCTG 44901

RESULT 8
US-09-949-016-16138
; Sequence 16138, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16138
; LENGTH: 126176
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-16138

Query Match 72.8%; Score 18.2; DB 4; Length 126176;
Best Local Similarity 87.0%; Pred. No. 1.8e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||

Db 44879 GAAGAAATTCAGTTCCTAGCTG 44901

RESULT 9

US-09-949-016-12387/c
; Sequence 12387, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12387
; LENGTH: 223471
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(223471)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-12387

Query Match 72.8%; Score 18.2; DB 4; Length 223471;
Best Local Similarity 87.0%; Pred. No. 1.9e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||

Db 222918 GAAGAAATTCAGTTCATGTTG 222896

RESULT 10
US-09-949-016-12724/c
; Sequence 12724, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12724
; LENGTH: 223471
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(223471)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-12724

Query Match 72.8%; Score 18.2; DB 4; Length 223471;
Best Local Similarity 87.0%; Pred. No. 1.9e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||

Db 222918 GAAGAAATTCAGTTCATGTTG 222896


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; APPLICANT: KIMMERLY, WILLIAM JOHN
; TITLE OF INVENTION: STAPHYLOCOCCUS EPIDERMIDIS NUCLEIC ACIDS AND PROTEINS
; FILE REFERENCE: PU3480US
; CURRENT APPLICATION NUMBER: US/09/710,279
; CURRENT FILING DATE: 2000-11-09
; PRIOR APPLICATION NUMBER: 60/164,258
; PRIOR FILING DATE: 1999-11-09
; NUMBER OF SEQ ID NOS: 4472
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 999
; LENGTH: 1325
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: synthetic
; OTHER INFORMATION: nucleic acid sequence
US-09-710-279-999

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Query Match      71.2%; Score 17.8; DB 4; Length 1325;
Best Local Similarity 90.5%; Pred. No. 1.6e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Qy      5 GAAATTCAGTTCATAGCTTGT 25
      |||||
Db      1305 GAAATTCAGTTCATAGCTTGT 1285

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Search completed: August 13, 2005, 06:48:53
Job time : 65.4362 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 13, 2005, 03:26:33 ; Search time 1606.65 Seconds
(without alignments)
592.293 Million cell updates/sec

Title: US-10-673-854-3

Perfect score: 25
Sequence: 1 tgaagaaattcagttcattgt 25

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST:*
1: gb_est1:*
2: gb_est2:*
3: gb_est3:*
4: gb_est4:*
5: gb_est5:*
6: gb_est6:*
7: gb_est7:*
8: gb_gss1:*
9: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20.2	80.8	187	6	CD025522 NXSI_060
2	20.2	80.8	623	7	CO198363 GRC1_13.D
3	20.2	80.8	780	7	CF689706 RUCNT1_45
4	20.2	80.8	902	7	CO226474 WS01029.B
5	19.8	79.2	280	5	BQ236799 TAE05026F
6	19.8	79.2	395	6	CA707706 wdk2c.pk0
7	19.8	79.2	546	6	CD895321 G174_001I
8	19.8	79.2	581	2	BE402372 CSB007C06
9	19.8	79.2	581	5	BQ607758 BRY_3653
10	19.8	79.2	634	4	BQ250598 BU250598
11	19.8	79.2	636	6	CD896013 G174_101J
12	19.8	79.2	643	6	CD453897 WHE0902.E
13	19.8	79.2	773	5	BQ804516 WHE3555.F
14	19.8	79.2	783	8	BZ186617 CH230_340
15	19.8	79.2	863	9	CC495993 CH240_331
16	19.8	79.2	1002	9	CNS05LGB AL342737 Tetrarodon
17	19.4	77.6	710	7	CK367386 AGENCOURT
18	19.2	76.8	506	9	CG639240 OST369798
19	19.2	76.8	536	7	CN518432 G00094.B3
20	19.2	76.8	588	8	AZ229308 RRC1-23-5
21	19.2	76.8	599	8	AQ987207 RRC1-23-3
22	19.2	76.8	662	5	BQ139159 NF011D06P
23	19.2	76.8	689	8	B92536 CIT-HSP-217
24	19.2	76.8	725	9	CE361033 tigr-gss-

C 25	19.2	76.8	775	7	CN986329	61938.125
C 26	19.2	76.8	810	5	BU333932	603499379
C 27	19.2	76.8	961	4	BG760956	60217652
C 28	19.2	76.8	963	9	CL053389	CH216-77J
C 29	19.2	76.8	1291	9	AG435478	Mus muscu
C 30	18.8	75.2	231	8	BZ416853	IF71N09.G
C 31	18.8	75.2	529	6	CD312483	StrPu691.
C 32	18.8	75.2	544	8	AZ007802	RFC1-23-3
C 33	18.8	75.2	619	4	BG085680	H3116C06-
C 34	18.8	75.2	699	8	AZ106414	RPC1-23-3
C 35	18.8	75.2	704	1	AL698618	DKFZp686K
C 36	18.8	75.2	726	9	CE132344	tigr-gss-
C 37	18.8	75.2	1058	9	CNS05MCC	AL343893 Tetrarodon
C 38	18.8	75.2	2625	9	CL960736	OSIFCC037
C 39	18.6	74.4	143	6	CD313427	StrPu621.
C 40	18.6	74.4	241	9	CL325373	RPC144.24
C 41	18.6	74.4	275	8	AQ103010	HS_3059.A
C 42	18.6	74.4	349	5	BQ32895	QV0-ET014
C 43	18.6	74.4	354	2	AW756746	SI26901.Y
C 44	18.6	74.4	383	8	AQ238508	RPC111-63
C 45	18.6	74.4	425	4	BM176564	TGESTzya9

ALIGNMENTS

RESULT 1
LOCUS CD025522 187 bp mRNA linear EST 07-MAY-2003
DEFINITION NXSI_060_F12_F NXSI (Nsf Xylem Side wood Inclined) Pinus taeda cDNA
clone NXSI_060_F12_5', mRNA sequence.
ACCESSION CD025522
VERSION CD025522.1 GI:30364172
KEYWORDS EST.
SOURCE Pinus taeda (loblolly pine)
ORGANISM Pinus taeda
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Coniferopsida; Coniferales; Pinaceae; Pinus; Pinus.
1 (bases 1 to 187)
AUTHORS Sederoff, R.
TITLE Molecular Basis of Wood Formation in the Pine Megagenome
JOURNAL Unpublished (2000)
COMMENT Contact: Sederoff, Ron
Forest Biotechnology
North Carolina State University
840 Main Campus Drive, Centennial Campus, Campus Box 7247, Raleigh,
NC 27695, USA
Tel: 919 515 7800
Fax: 919 515 7801
Email: ron.sederoff@ncsu.edu, jerri.johnson@ncsu.edu
Please see <http://web.abc.umn.edu/biodata/nsfpine/> for further
information.
Seq primer: T3.

FEATURES
Location/Qualifiers
1..187
/organism="Pinus taeda"
/mol_type="mRNA"
/strain="Coastal plain loblolly pine from North Carolina"
/db_xref="taxon:3352"
/clone="NXSI_060_F12"
/tissue_type="Xylem"
/cell_type="Side"
/dev_stage="Juvenile"
/lab_host="XL1-Blue"
/clone_lib="NXSI (Nsf Xylem Side wood Inclined)"
/note="Vector: Bluescript SK; Site 1: Eco RI; Site 2:
XhoI; The library is from early (spring) wood, taken from
three six-year old trees (three different genotypes), in
the juvenile phase. These trees were induced to form side
wood by bending to a 45 degree angle and tying them to the
ground. Differentiating xylem was harvested from the sides
of the inclined stems, and a mixture of all three
genotypes was used for the library. oligo-dT primed cDNA

was directionally cloned into the EcoRI-XhoI Bluescript SK vector arms. NOTE: The sequences contain a 'cDNA adapter' between the EcoRI site and the start of the EST. The adapter sequence is 'AATTCGGCAGG'.

ORIGIN
Query Match 80.8%; Score 20.2; DB 6; Length 187;
Best Local Similarity 88.0%; Pred. No. 2e+02;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTCATAGCTTGT 25
|||||
Db 125 TGATGAATTCAGTTCATACCATGT 149

RESULT 2
CO198363
LOCUS
DEFINITION
cDNA clone GEO1_13_D06_A029 3', mRNA sequence.
CO198363
VERSION
KEYWORDS
SOURCE
ORGANISM
Pinus taeda (loblolly pine)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Coniferales; Pinaceae; Pinus; Pinus.
REFERENCE
AUTHORS
Pratt,L., Cordonnier-Pratt,M.-M., Lorenz,W.W., Zimmermann,C. and Dean,J.F.D.
TITLE
An EST database from gravitropically stimulated loblolly pine (Pinus taeda) roots
JOURNAL
COMMENT
Other_ESTRs: GEO1_13_D06.g1_A029
Contact: Cordonnier-Pratt MM
Laboratory for Genomics and Bioinformatics
The University of Georgia, Department of Plant Biology
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
Tel: 706 542 1860
Fax: 706 583 0210
Email: mmpratt@uga.edu

RNA prepared and library constructed by W. Walter Lorenz (School of Forest Resources, University of Georgia); plant material prepared by Craig Zimmermann (School of Forest Resources, University of Georgia) using rooted cuttings provided by the Forest Biology Research Cooperative (FBRC) and the CCLONES project a the University of Florida; sequencing done in the Laboratory for Genomics and Bioinformatics, University of Georgia. Sequence ends have been trimmed to exclude vector and regions below Phred quality 16. Three-prime sequences are presented as their reverse complement and have been trimmed to exclude polyA.
Seq primer: M13-21 (TGTAACGACGCGCAGT)
POLYA=Yes.

FEATURES
source
1. .623
/organism="Pinus taeda"
/mol_type="mRNA"
/strain="3 CCLONES"
/db_xref="taxon:3352"
/clone="GEO1_13_D06_A029"
/lab_host="DH10B-T1 phage-resistant E. coli"
/note="Vector: pSL180; Site 1: EcoRI; Site2: XhoI; The library was prepared from polyA+ RNA from the roots of 1-year-old loblolly pine (Pinus taeda) cuttings that were rooted and then planted in washed sand. The rooted cuttings were maintained for 27 days (April 2003 harvest) under ambient conditions in a local greenhouse. They were kept on a weekly regimen of 0.5x nutrient-complete Hoagland's solution and supplemented with additional water sufficient to maintain a 15% soil moisture content. Twenty-four hours (24 h) prior to harvesting roots for mRNA preparation, the potted trees were tipped 90 degree

ORIGIN

Query Match 80.8%; Score 20.2; DB 7; Length 623;
Best Local Similarity 88.0%; Pred. No. 2.3e+02;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TGAAGAAATTCAGTTCATAGCTTGT 25
|||||
Db 180 TGATGAATTCAGTTCATACCATGT 204

RESULT 3
CF669706
LOCUS
DEFINITION
RTCN1_45_C06_g1_A029 Root control Pinus taeda cDNA clone
RTCN1_45_C06_A029 5', mRNA sequence.
CF669706
VERSION
KEYWORDS
SOURCE
ORGANISM
Pinus taeda (loblolly pine)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Coniferales; Pinaceae; Pinus; Pinus.
REFERENCE
AUTHORS
Pratt,L., Cordonnier-Pratt,M.-M., Lorenz,W.W., Zimmermann,C. and Dean,J.F.D.
TITLE
An EST database from untreated loblolly pine (Pinus taeda) roots
JOURNAL
COMMENT
Contact: Cordonnier-Pratt MM
Laboratory for Genomics and Bioinformatics
The University of Georgia, Department of Plant Biology
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
Tel: 706 542 1860
Fax: 706 583 0210
Email: mmpratt@uga.edu

RNA prepared and library constructed by W. Walter Lorenz (School of Forest Resources, University of Georgia); plant material prepared by Craig Zimmermann (School of Forest Resources, University of Georgia) using rooted cuttings provided by the Forest Biology Research Cooperative (FBRC) and the CCLONES project a the University of Florida; sequencing done in the Laboratory for Genomics and Bioinformatics, University of Georgia. Sequence ends have been trimmed to exclude vector and regions below Phred quality 16. Three-prime sequences are presented as their reverse complement and have been trimmed to exclude polyA.
Seq primer: JENREV (CAGGAACAGCTATGACC).

FEATURES
source

1. .780
/organism="Pinus taeda"
/mol_type="mRNA"
/strain="3 CCLONES"
/db_xref="taxon:3352"
/clone="RTCN1_45_C06_A029"
/lab_host="DH10B-T1 phage-resistant E. coli"
/clone_lib="Root control"
/note="Organ: root; Vector: pSL1180; Site 1: EcoRI; Site 2: XhoI; The library was prepared from polyA+ RNA from the roots of 1-year-old loblolly pine (Pinus taeda) cuttings that were rooted and then planted in washed sand. Just before harvesting roots for RNA isolation, the rooted cuttings were maintained for 27 days (April 2003) under ambient conditions in a local greenhouse. They were kept on a weekly regimen of 0.5x nutrient-complete Hoagland's solution and supplemented with additional water sufficient to maintain a 15% soil moisture content. Double-stranded cDNA was cloned unidirectionally into pSL180. Inserts can be excised with EcoRI (5' end) and XhoI (3' end)."

ORIGIN

ORIGIN

Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Poideae; Triticeae; Triticum.

1 (bases 1 to 395)
Tingey, S.V., Powell, W., Wolters, P., Dolan, M., Hainey, C., Yuan, Z.,
Miao, G., Caraher, N. and Hanafey, M.K.
DuPont Wheat cDNA Sequence
Unpublished (2002)

Contact: Scott V. Tingey
Crop Genetics
E. I. DuPont de Nemours and Company

1 Innovation Way, P.O. Box 6104, Newark, DE 19714-6104, USA
Tel: 302-631-2602
Fax: 302-631-2607
Email: Scott.V.Tingey@USA.dupont.com
Seq primer: M13.

FEATURES
source

Location/Qualifiers
1..395
/organism="Triticum aestivum"
/mol_type="mRNA"
/db_xref="taxon:4565"
/clone="wdk2c.pk006.b16"
/tissue_type="kernel"
/clone_lib="wdk2c"
/note="Vector: pBluescript SK+; Site 1: EcoRI; Site 2:
XhoI; Wheat (Triticum aestivum L.) developing kernel, 7
days after anthesis."

ORIGIN

Query Match 79.2%; Score 19.8; DB 6; Length 395;
Best Local Similarity 91.3%; Pred. No. 3.3e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||
DB 327 GAAGAAATTCAGTTCCTAGCATG 305

RESULT 7
CD895321/c
LOCUS
DEFINITION
G174.00115F010514 G174 Triticum aestivum cDNA clone G17400115,
mRNA sequence.

ACCESSION
CD895321.1 GI:32667779
VERSION
EST.
KEYWORDS
SOURCE
ORGANISM
Triticum aestivum (bread wheat)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Poideae; Triticeae; Triticum.

1 (bases 1 to 546)

Genoplante.
Genoplante, a major partnership french program in plant genomics
Unpublished (2003)
Contact: Genoplante

Genoplante
93, rue Henri Rochefort 91025 EVRY CEDEX France
Tel: 33 1 69 47 54 00
Fax: 33 1 69 47 54 10

This sequence has been generated in the framework of the french
plant genomics programme 'Genoplante' (<http://www.genoplante.com>
and <http://genoplante-info.infobiogen.fr>).

FEATURES
source

Location/Qualifiers
1..546
/organism="Triticum aestivum"
/mol_type="mRNA"
/cultivar="recital"
/db_xref="taxon:4565"
/clone="G17400115"
/tissue_type="grain (174 degrees per day after
pollination)"
/clone_lib="G174"

ORIGIN

1 (bases 1 to 546)
BQ607758
LOCUS
DEFINITION
BRY 3653 wheat EST endosperm library Triticum aestivum cDNA 5',
mRNA sequence.

Query Match 79.2%; Score 19.8; DB 6; Length 546;
Best Local Similarity 91.3%; Pred. No. 3.5e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||
DB 486 GAAGAAATTCAGTTCCTAGCATG 464

RESULT 8
BE402372/c
LOCUS

DEFINITION
CS8007C06F90908 ITEC CSB Wheat Endosperm Library Triticum aestivum
cDNA clone CSB007C06, mRNA sequence.

ACCESSION
BE402372
VERSION
KEYWORDS
SOURCE
ORGANISM
Triticum aestivum (bread wheat)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Poideae; Triticeae; Triticum.

1 (bases 1 to 581)

REFERENCE

AUTHORS
Anderson, O.A., Appels, R., Bailey, P., Blake, T., Close, T.,
Cloutier, S., Dubcovsky, J., Feuillet, C., Gale, M., Graner, A.,
Gustafson, P., Herrmann, R.G., Holton, T., Jacquemin, J.M., Jia, J.,
Joudrier, P., Langridge, P., Lazo, G.R., Lin, J.J., McGuire, P.,
Ogihara, Y., Pecchioni, N., Qualset, C., Schuch, W., Selvaraj, G.,
Shariflou, M., Sorrells, M., Warburton, M. and Wenzel, G.
International Triticeae EST Cooperative (ITEC): Production of
Expressed Sequence Tags for Species of the Triticeae
Unpublished (2000)
Contact: Appels R
Div. of Plant Industry, CSIRO
Canberra ACT 2601 AUSTRALIA
Tel: 61 62 465496
Fax: 61 62 485000
Email: rudi@pi.csiro.au
International Triticeae EST Cooperative (ITEC)
<http://wheat.pw.usda.gov/genome>.

FEATURES

source

Location/Qualifiers
1..581
/organism="Triticum aestivum"
/mol_type="mRNA"
/cultivar="Wyuna"
/db_xref="taxon:4565"
/clone="CSB007C06"
/tissue_type="endosperm"
/dev_stage="8-12 days post anthesis"
/lab_host="Escherichia coli SOLR"
/clone_lib="ITEC CSB Wheat Endosperm Library"
/note="Vector: Lambda Zap/Bluescript; Site 1: XhoI;
Site 2: EcoRI; Plants grown in Phytotron with 18C/13C
(day/night) 16 hour light. M13 Reverse sequencing primer
used. 1.0 Kbp average insert size."

ORIGIN

Query Match 79.2%; Score 19.8; DB 2; Length 581;
Best Local Similarity 91.3%; Pred. No. 3.5e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||
DB 575 GAAGAAATTCAGTTCCTAGCATG 553

RESULT 9

BQ607758/c
LOCUS
DEFINITION
BQ607758 wheat EST endosperm library Triticum aestivum cDNA 5',
mRNA sequence.

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VERSION BQ607758.1 GI:21557087
KEYWORDS EST.
SOURCE Triticum aestivum (bread wheat)
ORGANISM Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Poideae; Triticeae; Triticum.
REFERENCE 1 (bases 1 to 581)
AUTHORS Clarke, B., Lambrecht, M. and Rhee, S.Y.
TITLE Arabidopsis genomic information for interpreting wheat EST
sequences
JOURNAL Arabidopsis Integr. Genomics 3 (1-2), 33-38 (2003)
MEDLINE 22478026
PUBMED 12590341
COMMENT Contact: Lambrecht M
The Arabidopsis Information Resource
Carnegie Institution of Washington, Dept. of Plant Biology
260 Panama Street, Stanford, CA 94305, USA
Tel: 1 650 325 1521 x 251
Fax: 1 650 325 3748
Email: rhee@acoma.stanford.edu.
FEATURES
source
1..581
/organism="Triticum aestivum"
/mol_type="mRNA"
/cultivar="Wyuna"
/db_xref="taxon:4565"
/tissue_type="endosperm"
/dev_stage="developing endosperm tissue 8, 10 and 12 DPA
(days post anthesis)"
/clone_lib="wheat EST endosperm library"
ORIGIN
Query Match 79.2%; Score 19.8; DB 5; Length 581;
Best Local Similarity 91.3%; Pred. NO. 3.5e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||
Db 575 GAAGAAATTCAGTTCCTAGCATG 553

RESULT 10
BJ250598 634 bp mRNA linear EST 05-APR-2002
LOCUS BJ250598 Y. Ogiwara unpublished cDNA library, wh_f Triticum
DEFINITION aestivum cDNA clone wh16f11 3', mRNA sequence.
ACCESSION BJ250598
VERSION BJ250598.1 GI:20060583
KEYWORDS EST.
SOURCE Triticum aestivum (bread wheat)
ORGANISM Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Poideae; Triticeae; Triticum.
REFERENCE 1 (bases 1 to 634)
AUTHORS Ogiwara, Y. and Murai, K.
TITLE Expressed genes in Triticum aestivum
JOURNAL Unpublished (2002)
COMMENT Contact: Tadasu Shin-i
Center For Genetic Resource Information
National Institute of Genetics
1111 Yata, Mishima, Shizuoka 411-8540, Japan
Tel: 81-559-81-6856
Fax: 81-559-81-6855
Email: tshini@genes.nig.ac.jp.
FEATURES
source
1..634
/organism="Triticum aestivum"
/mol_type="mRNA"
/cultivar="Chinese Spring"
/db_xref="taxon:4565"
/clone="wh16f11"

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ORIGIN
Query Match 79.2%; Score 19.8; DB 4; Length 634;
Best Local Similarity 91.3%; Pred. NO. 3.5e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||
Db 29 GAAGAAATTCAGTTCCTAGCATG 51

RESULT 11
CD896013 636 bp mRNA linear EST 14-JUL-2003
LOCUS G174.101JL6R011120 G174 Triticum aestivum cDNA clone G174101J16,
DEFINITION mRNA sequence.
ACCESSION CD896013
VERSION CD896013.1 GI:32669222
KEYWORDS EST.
SOURCE Triticum aestivum (bread wheat)
ORGANISM Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Poideae; Triticeae; Triticum.
REFERENCE 1 (bases 1 to 636)
AUTHORS Genoplante.
TITLE Genoplante, a major partnership french program in plant genomics
JOURNAL Unpublished (2003)
COMMENT Contact: Genoplante
Genoplante
93, rue Henri Rochefort 91025 EVRY CEDEX France
Tel: 33 1 69 47 54 00
Fax: 33 1 69 47 54 10
This sequence has been generated in the framework of the french
plant genomics programme 'Genoplante' (http://www.genoplante.com
and http://genoplante-info.infobiogen.fr).
FEATURES
source
1..636
/organism="Triticum aestivum"
/mol_type="mRNA"
/cultivar="recital"
/db_xref="taxon:4565"
/clone="G174101J16"
/tissue_type="grain (174 degrees per day after
pollination)"
/clone_lib="G174"
ORIGIN
Query Match 79.2%; Score 19.8; DB 6; Length 636;
Best Local Similarity 91.3%; Pred. NO. 3.6e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 2 GAAGAAATTCAGTTCATAGCTTG 24
|||||
Db 69 GAAGAAATTCAGTTCCTAGCATG 91

RESULT 12
CD453897 643 bp mRNA linear EST 03-JUN-2003
LOCUS WHE0902_E02_I04ZT CS wheat 5-15 DAP spike cDNA library Triticum
DEFINITION aestivum cDNA clone WHE0902_E02_I04, mRNA sequence.
ACCESSION CD453897
VERSION CD453897.1 GI:31368525
KEYWORDS EST.
SOURCE Triticum aestivum (bread wheat)
ORGANISM Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Poideae; Triticeae; Triticum.

```

REFERENCE 1 (bases 1 to 643)
 AUTHORS Anderson,O.D., Butler,E., Chao,S., Choi,D.W., Close,T.J., Crossman,C., Fenton,R.D., Lazo,G.R., Pham,J., Rausch,C.J. and Woo,J.
 TITLE The structure and function of the expressed portion of the wheat
 JOURNAL genomes - 5-15 DAP CS spike cDNA library
 COMMENT Unpublished (2003)
 Contact: Olin Anderson
 US Department of Agriculture, Agriculture Research Service, Pacific
 West Area, Western Regional Research Center
 800 Buchanan Street, Albany, CA 94710, USA
 Tel: 5105595773
 Fax: 5105595818
 Email: oandern@pw.usda.gov
 This EST was generated by sequencing from the 3' end of the clone.
 Sequences have been trimmed to remove vector sequence and low
 quality sequence with phred score less than 20.
 Seq primer: T7 primer.

FEATURES
 source
 Location/Qualifiers
 1..643
 /organism="Triticum aestivum"
 /mol_type="mRNA"
 /cultivar="Chinese Spring"
 /db_xref="taxon:4565"
 /clone="WHE0902.E02.I04"
 /tissue_type="Spike"
 /dev_stage="Adult plant"
 /lab_host="E. coli SOLR"
 /note="Vector: Lambda Uni-ZAP XR, excised phagemid;
 Site 1: EcoRI; Site 2: XhoI; Plants were grown in the
 greenhouse. Spikes at 5, 10 and 15 DAP were harvested,
 total RNA and poly(A) RNA were prepared, a cDNA library
 was made, and the cDNA clones were in vivo excised to give
 phuescript phagemids in the TJ Close lab (Choi, Close,
 Fenton) at the University of California, Riverside.
 Plasmid DNA preparations and DNA sequencing were performed
 in the OD Anderson lab (all other authors)."

ORIGIN
 Query Match 79.2%; Score 19.8; DB 6; Length 643;
 Best Local Similarity 91.3%; Pred. No. 3.6e+02;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATGCTTG 24
 |||||
 DB 96 GAAGAAATTCAGTTCCTAGCATG 118
 |||||

RESULT 13
 BQ804516/c
 LOCUS
 DEFINITION WHE3555_F02_K03ZS Wheat developing grains cDNA library Triticum
 aestivum cDNA clone WHE3555_F02_K03, mRNA sequence.
 ACCESSION BQ804516
 VERSION BQ804516.1 GI:22028647
 KEYWORDS EST.
 SOURCE Triticum aestivum (bread wheat)
 ORGANISM Triticum aestivum
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Poideae; Triticeae; Triticum.
 1 (bases 1 to 773)
 Altenbach,S., Anderson,O.D., Chao,S., Chin,A., Close,T.J.,
 Cronin,K., Crossman,C., Fenton,R.D., Lazo,G.R., Pham,J.,
 Rausch,C.J., Wilson,C. and Woo,J.
 The structure and function of the expressed portion of the wheat
 genomes - Developing grains cDNA library
 Unpublished (2002)
 Contact: Olin Anderson
 US Department of Agriculture, Agriculture Research Service, Pacific
 West Area, Western Regional Research Center
 800 Buchanan Street, Albany, CA 94710, USA

Tel: 5105595773
 Fax: 5105595818
 Email: oandern@pw.usda.gov
 Sequences have been trimmed to remove vector sequence and low
 quality sequence with phred score less than 20
 Seq primer: SK primer.

FEATURES
 source
 Location/Qualifiers
 1..773
 /organism="Triticum aestivum"
 /mol_type="mRNA"
 /cultivar="Butte 86"
 /db_xref="taxon:4565"
 /clone="WHE3555_F02_K03"
 /tissue_type="whole grains"
 /dev_stage="3-44 days post anthesis seed"
 /lab_host="E. coli SOLR"
 /note="Vector: Lambda ZAP II, excised phagemid; Site 1:
 EcoRI; Plants were grown under six following different
 environmental regimes in greenhouse, Environment 1)
 240C/170C day/night, well-watered, with post-anthesis
 fertilizer, Environment 2) 240C/170C day/night,
 well-watered, without post-anthesis fertilizer,
 Environment 3) 370C/170C day/night, well-watered, with
 post-anthesis fertilizer, Environment 4) 370C/170C
 day/night, well-watered, without post-anthesis fertilizer,
 Environment 5) 370C/170C day/night plus drought, with
 post-anthesis fertilizer, Environment 6) 370C/170C
 day/night plus drought, without post-anthesis fertilizer,
 and developing wheat grains from the following were excised
 and frozen in liquid nitrogen, Environment 1 at 3, 5, 7,
 8, 10, 12, 16, 20, 24, 28, 32, 36, 40, 44 DPA Environment
 2 at 3, 5, 7, 8, 10, 12, 16, 20, 24, 28, 32, 36, 40, 44
 DPA Environment 3 at 3, 5, 7, 8, 10, 12, 16, 20, 24, 28,
 32, 34 DPA Environment 4 at 3, 5, 7, 8, 10, 12, 16, 20,
 24, 28, 32, 34 DPA Environment 5 at 3, 5, 7, 8, 10, 12,
 16, 20, 24, 28, 30 DPA Environment 6 at 3, 5, 7, 8, 10,
 12, 16, 20, 24, 28, 30 DPA and total RNA was prepared by
 S. Altenbach and K. Cronin at USDA-ARS, Albany, CA. A
 cDNA library was made using poly (A) RNA, and the cDNA
 clones were in vivo excised to give phuescript SK(-)
 phagemids in the TJ Close lab (Chin, Close, Fenton) at
 the University of California, Riverside. Plasmid DNA
 preparations and DNA sequencing were performed in the OD
 Anderson lab (others)."

ORIGIN
 Query Match 79.2%; Score 19.8; DB 5; Length 773;
 Best Local Similarity 91.3%; Pred. No. 3.6e+02;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAAGAAATTCAGTTCATGCTTG 24
 |||||
 DB 754 GAAGAAATTCAGTTCCTAGCATG 732
 |||||

RESULT 14
 BZ186617
 LOCUS
 DEFINITION CH230-340J18_TV CHORI-230 Segment 2 Rattus norvegicus genomic clone
 CH230-340J18, genomic survey sequence.
 ACCESSION BZ186617
 VERSION BZ186617.1 GI:23838548
 KEYWORDS GSS.
 SOURCE Rattus norvegicus (Norway rat)
 ORGANISM Rattus norvegicus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
 Rattus.
 1 (bases 1 to 783)
 Zhao,S., Shetty,J., Shatsman,S., Tsegaye,G., Geer,K.,
 Shvartsbeyn,A., Gebregeorgis,E., Overton,L., Russell,D.,
 Riggs,F., de Jong,P. and Fraser,C.M.

TITLE Rat BAC End Sequences from Library CHORI-230 MboI segment
JOURNAL Unpublished (1999)
COMMENT Other GSSs: CH230-340J18.TJ
 Contact: Shaying Zhao
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: szhao@tigr.org
 Clones are derived from the rat BAC library CHORI-230
 (http://www.chori.org/bacpac/rat230.htm). For BAC library
 availability, please contact Pieter de Jong (pdejong@mail.cho.org).
 Clones may be purchased from BACPAC Resources
 (http://www.chori.org/bacpac/or ering information.htm). BAC end
 page: http://www.tigr.org/tdb/bac_ends/rat/bac_end_intro.html
 Plate: 340 row: J column: 18
 Seq primer: T7
 Class: BAC ends.

FEATURES
 source Location/Qualifiers
 1..783
 /organism="Rattus norvegicus"
 /mol_type="genomic DNA"
 /strain="BN/SENhad/MCW"
 /db_xref="taxon:10116"
 /clone="CH230-340J18"
 /sex="Female"
 /cell_type="Brain"
 /clone_lib="CHORI-230 Segment 2"
 /note="Vector: PTARBAC1.3; Site 1: MboI; Site 2: MboI;
 CHORI-230 rat (BN/SENhad/MCW) BAC library produced by
 Pieter de Jong"

ORIGIN

Query Match 79.2%; Score 19.8; DB 8; Length 783;
 Best Local Similarity 91.3%; Pred. No. 3.6e+02;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 TGAAGAAATTCAGTTCATAGCTT 23
 |||||
 Db 405 TGGAGAAATTCAGTTCATAGCTT 427

RESULT 15
CC495993
LOCUS CH240_331H6.T7 CHORI-240 Bos taurus genomic clone CH240_331H6,
DEFINITION genomic survey sequence.
ACCESSION CC495993
VERSION CC495993.1 GI:31809966
KEYWORDS GSS.
SOURCE Bos taurus (cow)
ORGANISM Bos taurus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Rutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 Bovinae; Bos.

REFERENCE

AUTHORS Holt, R., Stott, J., Yang, G., Barber, S., Smailus, D., Prabhu, A.-L.,
 Tsai, M., Cloutier, A., Lee, D., Girn, N., Olson, T., Mayo, M.,
 Butterfield, Y., Kirkpatrick, R., Liu, J., Guin, R., Chan, A., Chiu, R.,
 Mathewson, C., Wye, N., Masson, A., Brown-John, M., Jones, S.,
 Schein, J., Marra, M., de Jong, P., McWilliam, S., Barris, W.,
 Dalrymple, B. P. and Tellam, R.

TITLE Bovine BAC End Sequences from Library CHORI-240, PLATES 294 to 398
JOURNAL Unpublished (2003)
COMMENT Other GSSs: CH240_331H6.TARBAC13P2
 Contact: Rob Holt
 Sequencing
 The British Columbia Cancer Agency Genome Science Centre
 600 W. 10th Ave, Vancouver, British Columbia, Canada V5Z 4E6
 Tel: 604-877-6085
 Fax: 604-877-6276
 Email: rholt@bcgsc.ca

Clones are derived from the bovine BAC library CHORI-240
 (http://www.chori.org/bacpac/bovine240.htm). For BAC library
 availability, please contact Pieter de Jong (pdejong@mail.cho.org).
 Clones may be purchased from BACPAC Resources
 (http://www.chori.org/bacpac/ordering information.htm). This work
 was undertaken as part of the International Bovine BAC Mapping
 Consortium (IBBMC) by CSIRO Livestock Industries, Australia and the
 British Columbia Genome Sciences Centre, Canada.
 Plate: 331 row: H column: 6
 Seq primer: T7
 Class: BAC ends.

FEATURES
 source Location/Qualifiers
 1..863
 /organism="Bos taurus"
 /mol_type="genomic DNA"
 /strain="breed: Hereford"
 /db_xref="taxon:9913"
 /clone="CH240_331H6"
 /sex="Male"
 /cell_type="Blood"
 /clone_lib="CHORI-240"
 /note="Vector: PTARBAC1.3; Site 1: MboI; Site 2: MboI;
 Hereford bull Li Domino 99375; CHORI-240 Bovine BAC
 library (Male) produced by Pieter de Jong"

ORIGIN

Query Match 79.2%; Score 19.8; DB 9; Length 863;
 Best Local Similarity 91.3%; Pred. No. 3.7e+02;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 TGAAGAAATTCAGTTCATAGCTT 23
 |||||
 Db 394 TGAAGAAATTCAGTTCATAGATT 416

Search completed: August 13, 2005, 06:45:05
 Job time : 1614.65 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 13, 2005, 01:08:33 ; Search time 725.234 Seconds
(without alignments)
1536.704 Million cell updates/sec

Title: US-10-673-854-4

Perfect score: 23

Sequence: 1 caggagatcctgagattatgtgg 23

Scoring table: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl:*

1: gb_ba:*

2: gb_hgt:*

3: gb_in:*

4: gb_om:*

5: gb_ov:*

6: gb_pat:*

7: gb_ph:*

8: gb_pl:*

9: gb_pr:*

10: gb_ro:*

11: gb_sts:*

12: gb_sy:*

13: gb_un:*

14: gb_vi:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	23	100.0	212280	9 HSB362E11	AL121873 Human DNA
C 2	22	95.7	142099	9 AC002509	AC002509 Homo sapi
C 3	22	95.7	176872	9 AC012078	AC012078 Homo sapi
C 4	20.4	88.7	39904	9 AL590003	AL590003 Human DNA
C 5	20.4	88.7	41502	9 AL513262	AL513262 Human DNA
C 6	20.4	88.7	70140	9 AL139820	AL139820 Human DNA
C 7	20.4	88.7	85000	9 HSA312686	AJ312686 Homo sapi
C 8	20.4	88.7	86719	9 HSJ906P16	AL079339 Human DNA
C 9	20.4	88.7	87817	9 AC079174	AC079174 Homo sapi
C 10	20.4	88.7	88665	2 HSA132411	AJ132411 Homo sapi
C 11	20.4	88.7	111026	2 AL157403	AL157403 Homo sapi
C 12	20.4	88.7	113956	9 HS74M1	AL035704 Human DNA
C 13	20.4	88.7	119012	9 AC106732	AC106732 Homo sapi
C 14	20.4	88.7	128915	9 AC002416	AC002416 Human Chr
C 15	20.4	88.7	139203	2 AC068311	AC068311 Homo sapi
C 16	20.4	88.7	152531	2 AP005855	AP005855 Homo sapi
C 17	20.4	88.7	156361	2 AC026174	AC026174 Homo sapi
C 18	20.4	88.7	158028	2 AC019001	AC019001 Homo sapi
C 19	20.4	88.7	159593	2 AC026302	AC026302 Homo sapi

20	20.4	88.7	160373	2	AC024548	AC024548 Homo sapi
C 21	20.4	88.7	162616	9	AC099539	AC099539 Homo sapi
C 22	20.4	88.7	165261	2	AC012659	AC012659 Homo sapi
C 23	20.4	88.7	175448	2	AC018651	AC018651 Homo sapi
C 24	20.4	88.7	175598	2	AC136198	AC136198 Papio anu
C 25	20.4	88.7	177253	9	AC092024	AC092024 Homo sapi
C 26	20.4	88.7	181877	2	AC149458	AC149458 Papio anu
C 27	20.4	88.7	193589	9	AC025370	AC025370 Homo sapi
C 28	20.4	88.7	196361	9	AC005386	AC005386 citb 57.1
C 29	20.4	88.7	200034	9	AC107620	AC107620 Homo sapi
C 30	20.4	88.7	203407	2	AC006174	AC006174 Homo sapi
C 31	20.4	88.7	204116	9	CNS00000A	AL049828 Human chr
C 32	20.4	88.7	208547	2	AC021570	AC021570 Homo sapi
C 33	20.4	88.7	347253	9	AF363578	AF363578 Homo sapi
C 34	19.8	86.1	22955	6	AX780888	AX780888 Sequence
C 35	19.8	86.1	22955	9	AY485310	AY485310 Homo sapi
C 36	19.8	86.1	80515	9	AC011992	AC011992 Homo sapi
C 37	19.8	86.1	105788	9	AC020926	AC020926 Homo sapi
C 38	19.8	86.1	110000	2	AC145312_2	AC145312_2
C 39	19.8	86.1	110000	2	AC024562_0	AC024562_0
C 40	19.8	86.1	110000	2	AC024562_1	AC024562_1
C 41	19.8	86.1	110000	2	AC024562_2	AC024562_2
C 42	19.8	86.1	117213	9	AC026779	AC026779 Homo sapi
C 43	19.8	86.1	132150	9	AC005586	AC005586 Homo sapi
C 44	19.8	86.1	132150	9	AC005586	AC005586 Homo sapi
C 45	19.8	86.1	132150	9	AC005586	AC005586 Homo sapi

ALIGNMENTS

RESULT 1
HSB362E11/c
LOCUS
DEFINITION
Human DNA sequence from clone RP13-362E11 on chromosome X. Contains a pseudogene similar to mouse GEG-154 and moquitto MRRG, a pseudogene similar to human MMS2 and chicken CROC-1B, ESTs, STSs and GSSs, complete sequence.

ACCESSION
AL121873
VERSION
AL121873.15 GI:8218071
KEYWORDS
HTG; CROC-1B; GEG-154; MMS2; MRRG.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens

REFERENCE
1 (bases 1 to 212280)
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

AUTHORS
Wilson,S.
TITLE
Submitted (19-MAY-2000) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
JOURNAL
requests: clonerequest@sanger.ac.uk

COMMENT
On Jun 3, 2000 this sequence version replaced gi:7329905.

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em:, EMBL; Sw:, SWISSPROT; Tr:, TrEMBL; Wp:, WormPEP; Information on the WormPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/ChrX>

RP13-32E11 is from the library RP1-13.2 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://bacpac.med.buffalo.edu/VECTOR/pBACE3.6>

FEATURES

Source

```
1. .212280  
/organism="Homo sapiens"  
/mol_type="genomic DNA"  
/db_xref="taxon:9606"  
/chromosome="X"  
/clone="RP13-362E11"  
/clone_lib="PBC1-13"
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repeat_region	5. .42	/notes="MER34 repeat: matches 498. .534 of consensus"
repeat_region	120. .403	/notes="MER39b repeat: matches 68. .453 of consensus"
repeat_region	404. .498	/notes="MER39 repeat: matches 7. .103 of consensus"
repeat_region	502. .1573	/notes="LM4 repeat: matches 4008. .5095 of consensus"
repeat_region	1569. .1914	/notes="LM4 repeat: matches 3525. .3878 of consensus"
repeat_region	1906. .2066	/notes="LM6 repeat: matches 1376. .1528 of consensus"
repeat_region	2112. .2378	/notes="ALuJb repeat: matches 5. .274 of consensus"
repeat_region	2379. .2635	/notes="LM6 repeat: matches 1126. .1402 of consensus"
repeat_region	2651. .2697	/notes="LM2D repeat: matches 504. .550 of consensus"
repeat_region	2840. .3252	/notes="LM2D repeat: matches 77. .514 of consensus"
repeat_region	3351. .3420	/notes="LM2D repeat: matches 1. .71 of consensus"
repeat_region	3421. .3902	/notes="HERVL repeat: matches 5231. .5757 of consensus"
repeat_region	3903. .4462	/notes="280 copies 2 mer tt 57% conserved"
repeat_region	4465. .4782	/notes="ALuJb repeat: matches 1. .312 of consensus"
repeat_region	4783. .6107	/notes="HERVL repeat: matches 3896. .5242 of consensus"
repeat_region	6238. .6378	/notes="LM6 repeat: matches 973. .1115 of consensus"
repeat_region	6640. .6783	/notes="LMCS repeat: matches 7751. .7910 of consensus"
repeat_region	7282. .7567	/notes="LM2H repeat: matches 1. .302 of consensus"
misc_feature		complement(7766. .8138)
misc_feature		/note="match: GSS: Em:AQ098528"
misc_feature		complement(7769. .8123)
misc_feature		/note="match: GSS: Em:AQ105610"
misc_feature		complement(7779. .8205)
repeat_region		/note="match: GSS: Em:AQ239356"
repeat_region	8002. .8469	/note="LM2D repeat: matches 1. .503 of consensus"
repeat_region	8477. .8654	/notes="MIR repeat: matches 75. .256 of consensus"
repeat_region	8715. .8812	/note="49 copies 2 mer aa 61% conserved"
gene		complement(9391. .9893)
		/gene="BB362E11.1"
		/pseudo
CDS		complement(9391. .9893)
		/gene="BB362E11.1"
		/note="BB362E11.1 (pseudogene similar to mouse GEG-15)
		mosquito MRRG)
		match: CDNAS: Em:AK001774
		match: proteins: Sw:P50636 Tr:Q9Y4Y1 Tr:P90662 Tr:P91
		/pseudo
		/codon_start=1
		/evidence=not experimental

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misc_feature /note="match: STS: Em:AU047022"
              complement(26000..26305)
misc_feature /note="match: GSS: Em:AQ123830"
              26005..26229
misc_feature /note="match: GSS: Em:AZ007683"
              26009..26160
misc_feature /note="match: STS: Em:HSB017WB9"
              26009..26147
misc_feature /note="match: GSS: Em:AZ069301"
              26037..26181
misc_feature /note="match: GSS: Em:AQ871738"
              complement(26048..26369)
misc_feature /note="match: GSS: Em:AQ871742"
              26046..26360

```

```

Query Match 100.0%; Score 23; DB 9; Length 212280;
Best Local Similarity 100.0%; Pred.No.0.38;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Qy 1 CAGGAGATCTCGAGATTATGTGG 23
    |||||
Db 116939 CAGGAGATCTCGAGATTATGTGG 116917

```

```

RESULT 2
AC002509/c 142099 bp DNA linear PRI 25-NOV-1998
LOCUS Homo sapiens chromosome Y, clone 2Y, complete sequence.
DEFINITION AC002509
ACCESSION AC002509.1 GI:3927860
VERSION HTG.
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 142099)
AUTHORS Birren,B., Linton,L., Nusbaum,C. and Lander,B.
TITLE Homo sapiens chromosome Y, clone 2Y
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 142099)
AUTHORS Hawkins,T.L., Birren,B.W., Pasman,K.H., Nusbaum,C., Lander,B.S.,
          McKernan,K., Munro,C., Richardson,P., Barna,N., Chang,A., Cooke,P.,
          Daly,M.J., Devon,K., Dewar,K., Forrest,C., Gage,D., Geraghty,K.,
          Hagos,B., Huang,J., Hui,L., Jacotot,L., Kirby,A., Lane,M.,
          Mackenzie,J., Marquis,N., McDermott,J., Molla,M., Morrow,J.,
          Nachman,A., Naylor,J., Nusbaum,C., O'Connor,T., Olotu,A.,
          Peterson,K., Reeve,M.P., Roberts,D., Rollins,G., Stilwell,J.,
          Stone,C., Strickland,C., Sydney,K., Tang,L., Wilmer,F., Zemtseva,I.
          and Zody,M.

```

```

Direct Submission
TITLE Submitted (27-AUG-1997) Whitehead Institute/MIT Center for Genome
JOURNAL Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE 3 (bases 1 to 142099)
AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,B., Allen,N., Anderson,M.,
          Baker,J., Baldwin,J., Barna,N., Beckler,R., Benn,J., Boutwell,C.,
          Brown,A., Castle,A., Cerny,J., Colangelo,M., Collins,S.,
          Collymore,A., Cooke,P., Corliss,D., Depayre,E., Devon,K., Dewar,K.,
          Donelan,L., Ferreira,P., FitzHugh,W., Forrest,C., Funke,R.,
          Gage,D., Gardyna,S., Geraghty,K., Grant,G., Hagos,B., Heaford,A.,
          Herena,L., Horton,L., Howland,J.C., Jacotot,L., Jones,C., Kann,L.,
          Karatas,A., Lehoczy,J., MacDonald,P., Marquis,N., McEwan,P.,
          McGurk,A., McKernan,K., Meldrim,J., Molla,M., Morris,W., Morrow,J.,
          Mychaleckyj,J., Nahf,R., Naylor,J., Niloff,M., O'Connor,T., Roy,A.,
          O'Donnell,P., Pavlin,B., Peterson,K., Riley,R., Roberts,D., Roy,A.,
          Severy,P., Stange-Thomann,N., Stilwell,J., Stojanovic,N., Stone,C.,
          Subramanian,A., Tesfaye,S., Tichovolsky,N., Torruella-Miller,I.,
          Vassiliev,H., Vo,A., Wagner,A., Wheeler,J., Wu,Y., Wyman,D.,
          Ye,W.J., Zhao,J. and Zody,M.
Direct Submission
TITLE Submitted (25-NOV-1998) Whitehead Institute/MIT Center for Genome
JOURNAL Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT On Nov 25, 1998 this sequence version replaced gi:3924665.
All repeats were identified using RepeatMasker: Smit, A.F.A. &

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```

Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html.
FEATURES
Location/Qualifiers
Source
1..142099
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="Y"
/maps="Y"
/clone="2Y"
/clone_lib="unknown"
complement(315..438)
/rpt_family="MER61-internal"
complement(2160..2521)
/rpt_family="MER50"
2522..2578
/rpt_family="AluSg/x"
2859..2919
/rpt_family="MERS1-internal"
complement(2920..2978)
/rpt_family="(CA)n"
2990..3699
/rpt_family="MER4-internal"
complement(3700..3814)
/rpt_family="LTR8"
3815..4468
/rpt_family="MER4-internal"
4523..7011
/rpt_family="MER4-internal"
complement(7295..7598)
/rpt_family="AluSp"
9162..9349
/rpt_family="MER4-internal"
9361..9509
/rpt_family="MER4A2"
9510..9811
/rpt_family="MER4A"
complement(9812..9962)
/rpt_family="LIM1"
complement(10212..10830)
/rpt_family="LIM4"
complement(10831..11120)
/rpt_family="AluY"
complement(11121..11337)
/rpt_family="LIM4"
complement(11508..12117)
/rpt_family="LIMBc"
complement(12315..12626)
/rpt_family="LIMB"
complement(12923..13136)
/rpt_family="MIR"
complement(13246..13328)
/note="single-stranded terminator coverage."
13761..13782
/rpt_family="AT_rich"
14930..14950
/rpt_family="AT_rich"
15128..15231
/rpt_family="HAL1"
15292..15552
/rpt_family="HAL1"
complement(15690..16275)
/rpt_family="MER4B"
complement(16397..16551)
/rpt_family="MER66-internal"
16825..16846
/rpt_family="AT_rich"
complement(17272..17680)
/rpt_family="MER4-internal"
complement(17687..17920)
/rpt_family="MER31-internal"
complement(18046..18789)
/rpt_family="MER50"

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repeat_region complement(19887..20110)
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repeat_region complement(20133..20299)
repeat_region /rpt_family="MER4-internal"
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repeat_region /rpt_family="AluY"
repeat_region complement(20594..20673)
repeat_region /rpt_family="MER4-internal"
repeat_region complement(20674..20974)
repeat_region /rpt_family="AluSg"
repeat_region complement(20975..21867)
repeat_region /rpt_family="MER4-internal"
repeat_region complement(21372..21498)
repeat_region /rpt_family="MER51-internal"
repeat_region complement(21499..21804)
repeat_region /rpt_family="AluSg"
repeat_region complement(21805..21964)
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repeat_region complement(22490..22671)
repeat_region /rpt_family="MER83-internal"
repeat_region complement(22589..22693)
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repeat_region 23324..23353
repeat_region /rpt_family="AT_rich"
repeat_region complement(23437..23977)
repeat_region /rpt_family="MER4B"
repeat_region complement(23978..24177)
repeat_region /rpt_family="MLT2A"
repeat_region 24557..24891
repeat_region /rpt_family="HAL1"
repeat_region 25476..25758
repeat_region /rpt_family="L2"
repeat_region 26069..26101
repeat_region /rpt_family="AT_rich"
repeat_region 26265..26489
repeat_region /rpt_family="MLT1D"
repeat_region 26490..26517
repeat_region /rpt_family="CAAA)n"
repeat_region 26518..26780
repeat_region /rpt_family="MLT1D"
repeat_region 27156..27180
repeat_region /rpt_family="AT_rich"
repeat_region complement(27417..27797)
repeat_region /rpt_family="L2"
repeat_region complement(27963..28332)
repeat_region /rpt_family="MSTA"
repeat_region complement(28333..29268)
repeat_region /rpt_family="MSTA-internal"
repeat_region 29269..29580
repeat_region /rpt_family="AluY"
repeat_region complement(29581..30301)
repeat_region /rpt_family="MSTA-internal"
repeat_region complement(30304..30679)
repeat_region /rpt_family="MSTA"
repeat_region complement(30680..31159)
repeat_region /rpt_family="LTR40a"
repeat_region complement(31325..31655)
repeat_region /rpt_family="LINE"
repeat_region 31692..31731
repeat_region /rpt_family="AT_rich"
repeat_region 31883..31927
repeat_region /rpt_family="AT_rich"
repeat_region complement(32107..33826)
repeat_region /rpt_family="LIM4"
repeat_region 33831..34558
repeat_region /rpt_family="LINE3"
repeat_region 34987..35012
repeat_region /rpt_family="AT_rich"
repeat_region 35108..35669

/rpt_family="MER4B"
36280..36322
/rpt_family="AT_rich"
complement(36386..36519)
/rpt_family="MIR"
complement(36755..37203)
/rpt_family="MER4C"

Query Match 95.7%; Score 22; DB 9; Length 142099;
Best Local Similarity 100.0%; Pred. No. 1.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 35610 CAGGAGATCCTGAGATTATGTG 35589

RESULT 3
AC012078/c 176872 bp DNA linear PRI 30-SEP-2000
LOCUS Homo sapiens BAC clone RP11-539022 from Y, complete sequence.
DEFINITION AC012078
ACCESSION AC012078
VERSION AC012078.3 GI:7684580
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 176872)
AUTHORS Sulston,J.E. and Waterston,R.
TITLE Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE 99063792
PUBMED 9847074
REFERENCE 2 (bases 1 to 176872)
AUTHORS Hou,S., Maupin,R. and Gibson,A.
TITLE The sequence of Homo sapiens BAC clone RP11-539022
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 176872)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (19-OCT-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE 4 (bases 1 to 176872)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (03-MAY-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE 5 (bases 1 to 176872)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (12-JUN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE 6 (bases 1 to 176872)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (30-SEP-2000) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On May 3, 2000 this sequence version replaced gi:7631053.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@watson.wustl.edu
----- Summary Statistics
----- Center project name: H_NH0539022

NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
```

clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

The position of this clone was established as part of a collaboration between the Human Chromosome Y Mapping Project (Tomoko Kawaguchi, Helen Skaletsky, Laura G. Brown, Steve Rozen, and David C. Page at the Whitehead Institute for Biomedical Research, Cambridge MA) and the Washington University Genome Sequencing Center, St. Louis MO.

SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Moon, P. Y., Zhao, B., Frengen, E., Tatenno, M., Catanesi, J. J., and de Jong, P. J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-530K5; the clone sequenced to the right is RP11-33605. Actual start of this clone is at base position 1 of RP11-539022; actual end is at base position 176872 of RP11-539022.

FEATURES

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  /mol_type="genomic DNA"
  /db_xref="taxon:9606"
  /chromosome="Y"
  /map="Y"
  /clone="RP11-539022"
  /clone_lib="RPCI-11"
repeat_region
1. .204
  /rpt_family="L1"
repeat_region
202. .1293
  /rpt_family="L1"
repeat_region
1294. .1324
  /rpt_family="L1"
repeat_region
1325. .2158
  /rpt_family="L1"
repeat_region
2159. .2339
  /rpt_family="L1"
repeat_region
2340. .3099
  /rpt_family="L1"
repeat_region
3120. .3219
  /rpt_family="Alu"
repeat_region
3220. .3945
  /rpt_family="L1"
repeat_region
3945. .4535
  /rpt_family="L1"
repeat_region
4565. .4619
  /rpt_family="Alu"
repeat_region
4638. .5206
  /rpt_family="L1"
repeat_region
5255. .5268
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repeat_region
5269. .5291
  /rpt_family="(TATG)n"
repeat_region
5292. .5583
  /rpt_family="Alu"
repeat_region
5989. .6021
  /rpt_family="AT_rich"

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6248. .6268
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repeat_region
6600. .6686
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repeat_region
7754. .8717
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8711. .10078
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10079. .10455
  /rpt_family="MaLR"
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10456. .11631
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repeat_region
11657. .11760
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11761. .12278
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12282. .12941
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12940. .12962
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12942. .13243
  /rpt_family="Alu"
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13241. .14024
  /rpt_family="L1"
repeat_region
14025. .14071
  /rpt_family="L1"
repeat_region
14072. .14107
  /rpt_family="(TAA)n"
repeat_region
14108. .14764
  /rpt_family="L1"
repeat_region
14770. .14800
  /rpt_family="AT_rich"
repeat_region
15566. .15598
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repeat_region
15847. .15867
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repeat_region
16123. .16431
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repeat_region
17318. .17558
  /rpt_family="L1"
repeat_region
17655. .18325
  /rpt_family="L1"
repeat_region
18319. .18728
  /rpt_family="L1"
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18735. .18866
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repeat_region
18880. .19147
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19173. .20018
  /rpt_family="L1"
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20015. .20440
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20441. .20536
  /rpt_family="LTR54"
repeat_region
20537. .21186
  /rpt_family="MER4-group"
repeat_region
21198. .21580
  /rpt_family="L1"
repeat_region
21581. .21618
  /rpt_family="(TTTA)n"
repeat_region
21671. .21838
  /rpt_family="L1"
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21839. .22135
  /rpt_family="Alu"
repeat_region
22136. .22315
  /rpt_family="L1"
repeat_region
24191. .24210
  /rpt_family="(TTG)n"
repeat_region
24211. .24482
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24828. .25925
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repeat_region
25914. .26164

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 repeat_region /note="MT1E repeat: matches 257. .392 of consensus" 31647. .31768
 repeat_region /note="MUT1D repeat: matches 326. .462 of consensus" 32760. .33055
 repeat_region /note="MT1J repeat: matches 214. .505 of consensus" 33110. .33413
 repeat_region /note="AluSg repeat: matches 1. .310 of consensus" 35009. .35548
 misc_feature /note="match: GSS: Em:AQ602626" 36014. .36132
 repeat_region /note="MER86 repeat: matches 19. .149 of consensus" 36608. .36657
 repeat_region /note="L1PB3 repeat: matches 6101. .6150 of consensus" 37258. .39904
 repeat_region /note="L1MAL repeat: matches 3664. .6297 of consensus"

ORIGIN

Query Match 88.7%; Score 20.4; DB 9; Length 39504;
 Best Local Similarity 95.5%; Pred. No. 9.8;
 Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22

Db 18372 CAGGAGATCCTGAGATTATGTG 18393

RESULT 5

AL513262
 LOCUS Human DNA sequence from clone RP11-499115 on chromosome 1, complete sequence.
 DEFINITION

ACCESSION

AL513262

VERSION

AL513262.7 GI:17381349

KEYWORDS

HTG.

SOURCE

Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 41502)

REFERENCE

Almeida, J.

Direct Submission

Submitted (30-NOV-2001) Wellcome Trust Sanger Institute, Hinxton,

Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk

On Dec 5, 2001 this sequence version replaced gi:16944082.

During sequence assembly data is compared from overlapping clones.

Where differences are found these are annotated as variations

together with a note of the overlapping clone name. Note that the

variation annotation may not be found in the sequence submission

corresponding to the overlapping clone, as we submit sequences with

only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all

regions were either double-stranded or sequenced with an alternate

chemistry or covered by high quality data (i.e., phred quality >=

30); an attempt was made to resolve all sequencing problems, such

as compressions and repeats; all regions were covered by at least

one plasmid subclone or more than one M13 subclone; and the

assembly was confirmed by restriction digest. The following

abbreviations are used to associate primary accession numbers given

in the feature table with their source databases: Em., EMBL; SW.,

SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information on the WORMPEP

database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormep/ This sequence

was generated from part of bacterial clone contigs of human

chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping

Group. Further information can be found at

http://www.sanger.ac.uk/HGP/Chr1

RP11-499115 is from the library RPC1-11.2 constructed by the group

of Pieter de Jong. For further details see

http://www.chori.org/bacpac/home.htm

VECTOR: pBACE3.6

IMPORTANT: This sequence is not the entire insert of clone

RP11-499115 It may be shorter because we sequence overlapping

sections only once, except for a short overlap.

The true left end of clone RP11-54388 is at 39503 in this sequence.

The true right end of clone RP4-553F17 is at 2000 in this sequence.

FEATURES

Location/Qualifiers

source 1. .41502

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/chromosome="1"

/clone="RP11-499115"

/clone_lib="RPC1-11.2"

ORIGIN

Query Match 88.7%; Score 20.4; DB 9; Length 41502;
 Best Local Similarity 95.5%; Pred. No. 9.8;
 Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22

Db 37869 CAGGAGATCCTGAGATTATGTG 37890

RESULT 6

AL139820
 LOCUS Human DNA sequence from clone RP11-396A24 on chromosome 10, complete sequence.
 DEFINITION

ACCESSION

AL139820

VERSION

AL139820.10 GI:14272240

KEYWORDS

HTG.

SOURCE

Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 70140)

REFERENCE

Ramsay, H.

Direct Submission

Submitted (30-MAY-2001) Sanger Centre, Hinxton, Cambridgeshire,

CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk

requests: clonerequest@sanger.ac.uk

On May 31, 2001 this sequence version replaced gi:13751278.

During sequence assembly data is compared from overlapping clones.

Where differences are found these are annotated as variations

together with a note of the overlapping clone name. Note that the

variation annotation may not be found in the sequence submission

corresponding to the overlapping clone, as we submit sequences with

only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all

regions were either double-stranded or sequenced with an alternate

chemistry or covered by high quality data (i.e., phred quality >=

30); an attempt was made to resolve all sequencing problems, such

as compressions and repeats; all regions were covered by at least

one plasmid subclone or more than one M13 subclone; and the

assembly was confirmed by restriction digest. The following

abbreviations are used to associate primary accession numbers given

in the feature table with their source databases: Em., EMBL; SW.,

SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information on the WORMPEP

database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormep/ This sequence

was generated from part of bacterial clone contigs of human

chromosome 10, constructed by the Sanger Centre Chromosome 10

Mapping Group. Further information can be found at

http://www.sanger.ac.uk/HGP/Chr10

RP11-396A24 is from the library RPC1-11.2 constructed by the group

of Pieter de Jong. For further details see

http://www.chori.org/bacpac/home.htm

VECTOR: pBACE3.6

IMPORTANT: This sequence is not the entire insert of clone

RP11-396A24 It may be shorter because we sequence overlapping

sections only once, except for a 100 base overlap.
The true right end of clone RP11-396A24 is at 70140 in this sequence. The true left end of clone RP11-343D22 is at 5760 in this sequence. The true right end of clone RP11-168C9 is at 100 in this sequence.

FEATURES

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/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="10"
/clone="RP11-396A24"
/clone_lib="RPC1-11.2"
1. .315
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324. .1989
/note="HERVL repeat: matches 4026. .5750 of consensus"
2022. .2050
/note="HERVL repeat: matches 3445. .3473 of consensus"
complement(2274. .2851)
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2847. .3318
/note="match: GSS: Em:AQ567074"
2972. .3102
/note="L1 repeat: matches 2572. .2750 of consensus"
3473. .3683
/note="L1 repeat: matches 40. .260 of consensus"
3887. .3998
/note="MIR repeat: matches 59. .178 of consensus"
4502. .4659
/note="MER5A repeat: matches 2. .170 of consensus"
4671. .4795
/note="MER81 repeat: matches 1. .114 of consensus"
5762. .6007
/note="match: GSS: Em:AQ395191"
5793. .6215
/note="MLT12 repeat: matches 1. .405 of consensus"
6322. .6496
/note="match: GSS: Em:AQ815465"
6338. .6628
/note="match: GSS: Em:A2517696"
6370. .6938
/note="MER4B repeat: matches 1. .575 of consensus"
8625. .8926
/note="AluY repeat: matches 1. .302 of consensus"
complement(8707. .9267)
/note="match: GSS: Em:AQ307791"
9958. .10246
/note="AluSg repeat: matches 1. .289 of consensus"
complement(11298. .11808)
/note="match: STS: Em:G49507"
complement(11361. .11848)
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11868. .12312
/note="match: GSS: Em:AQ369493"
13351. .13454
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13514. .13823
/note="AluJo repeat: matches 6. .310 of consensus"
14996. .15576
/note="match: STS: Em:G58481
match: GSS: Em:AQ347684"
18117. .18286
/note="5 copies 34 mer 74% conserved"
18122. .18277
/note="3 copies 52 mer 78% conserved"
18123. .18278
/note="78 copies 2 mer ca 76% conserved"
complement(19440. .19834)
/note="match: GSS: Em:AQ12568"
22796. .22835
/note="20 copies 2 mer ct 87% conserved"
22803. .22838

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23711. .23836
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27070. .27599
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27254. .27421
/note="FRAM repeat: matches -1. .168 of consensus"
27781. .28247
/note="HAL1 repeat: matches 419. .886 of consensus"
28623. .28757
/note="MIR repeat: matches 106. .243 of consensus"
28916. .29598
/note="match: GSS: Em:AQ390359"
29470. .30077
/note="L1MB2 repeat: matches 5551. .6167 of consensus"
30434. .30944
/note="L1MB3 repeat: matches 5375. .5901 of consensus"
31038. .31341
/note="AluSx repeat: matches 1. .304 of consensus"
31427. .31906
/note="L1MB repeat: matches 5226. .5728 of consensus"
32282. .32686
/note="L1MB4 repeat: matches 3515. .3937 of consensus"
32818. .33025
/note="L1MC5 repeat: matches 7355. .7555 of consensus"
33210. .33249
/note="20 copies 2 mer tt 85% conserved"
complement(33442. .33831)
/note="match: GSS: Em:B52135"
34349. .34584
/note="match: GSS: Em:AQ416077"
34611. .34920
/note="MLT1D repeat: matches 191. .533 of consensus"
36001. .36065
/note="L1MB repeat: matches 209. .282 of consensus"
36300. .36400
/note="L1 repeat: matches 2607. .2710 of consensus"
37143. .37455
/note="AluSx repeat: matches 1. .312 of consensus"
complement(37724. .38564)
/note="match: GSS: Em:AQ738968"
37958. .38287
/note="MLT1C repeat: matches 144. .466 of consensus"
38288. .38576
/note="AluJo repeat: matches 13. .297 of consensus"
38570. .39096
/note="match: GSS: Em:AQ337467"
38577. .38682
/note="MLT1C repeat: matches 38. .144 of consensus"
41198. .41284
/note="MIR repeat: matches 81. .173 of consensus"
41532. .41874
/note="MER47A repeat: matches 1. .366 of consensus"
41878. .41975
/note="MIR repeat: matches 59. .164 of consensus"
42637. .42726
/note="MIR repeat: matches 71. .164 of consensus"
complement(44297. .44694)
/note="match: GSS: Em:AQ545169"
45790. .46084
/note="AluSg repeat: matches 6. .297 of consensus"
47341. .47673
/note="MLT2E repeat: matches 1. .352 of consensus"
47675. .47722
/note="24 copies 2 mer ag 81% conserved"
47687. .47722
/note="6 copies 6 mer agagag 88% conserved"
complement(48625. .49183)
/note="match: GSS: Em:AQ556578"
49192. .49631
/note="match: GSS: Em:AQ675010"

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repeat_region 49940..50023
/notes="14 copies 6 mer ccacat 70% conserved"
repeat_region 50023..50323
/notes="AluSp repeat: matches 2..303 of consensus"
repeat_region 52327..52616
/notes="AluX repeat: matches 1..300 of consensus"
repeat_region 54026..54434
/notes="MLT2FB repeat: matches 4..408 of consensus"
repeat_region 54456..54874

Query Match      88.7%; Score 20.4; DB 9; Length 70140;
Best Local Similarity 95.5%; Pred. No. 9.8;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 6426 CAGGAGATCCTGAGATTATGTG 6447

RESULT 7
HSA312686      85000 bp DNA linear PRI 16-MAY-2002
LOCUS
DEFINITION Homo sapiens partial LIMD1 gene for LIM domains containing 1, exons
1-2, complete sequence.
ACCESSION AJ312686
VERSION AJ312686.1 GI:13548631
KEYWORDS HTG; LIM domains containing 1; LIMD1 gene.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Kiss,H., Yang,Y., Kiss,C., Andersson,K., Klein,G., Imreh,S. and
Dumanski,J.P.
TITLE The transcriptional map of the common eliminated region 1 (C3CER1)
in 3p21.3
JOURNAL Eur. J. Hum. Genet. 10 (1), 52-61 (2002)
MEDLINE 21906202
PUBMED 11896456
REFERENCE 2 (bases 1 to 85000)
AUTHORS Kiss,H.
TITLE Direct Submission
JOURNAL Submitted (01-APR-2001) Kiss H., Microbiology and Tumorbiology
Center (MTC), Karolinska Institute, Box 280, Stockholm, S-17177,
SWEDEN
COMMENT On Apr 9, 2001 this sequence version replaced gi:6599071.
FEATURES
source 1..85000
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="3"
/map="3p21.3"
/clone="RP5-1033n4"
24287..65692
/genes="LIMD1"
24287..25715
/genes="LIMD1"
/number=1
join(24308..25715,65591..65692)
/genes="LIMD1"
/codon_start=1
/product="LIM domains containing 1"
/protein_id="CAC35917.1"
/db_xref="GI:13548632"
/db_xref="GOA:Q9UGP4"
/db_xref="UniProt/TREMBL:Q9UGP4"
/translation="MDKYDDGLGEASKFTEDLMVYASDKGLFRVVDKAGNNPEPBT
RRVFATMAKHLQCCQQLQOETLPRGSRPVNGGRLGQARWEVVGSKLTVDGA
AKPPLAATGAPCAVTLLAQGPYPQQRSPRYLCTGRHSQDCGSRSLATSEMS
AFHQPGCDSPDCTGHDYDNLUSLASPKWGDKPGVPSIGLSVGWSPSSGSDPL
PKPCGDHPLNHLQSLSSSESGSLGSGNSSEKPTGLWSTASSQVRVSPGLP
SPNLENGAPAVGPVQPTPTSVSAPLALSCPRQGLPRNSGLGSEVGVMSKPNVDPQ

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PMFQDPKSYLSSAPSSPAGLDGSGQAVFGLGPKPGCTDLGTGPKLSPTSLVHPV
MSTLPELSCKEGLGWSGSLGSLVLDSPSPRVRLPCQPLVPGLPSPSAELKLE
AUTQRLEREMDAHPKADYFGACVKCKSGVFGAGQACQAMGNLYHDTCTCAAC"
25716..65590
/genes="LIMD1"
/number=1
65591..65692
/genes="LIMD1"
/number=2

Query Match      88.7%; Score 20.4; DB 9; Length 85000;
Best Local Similarity 95.5%; Pred. No. 9.7;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 17729 CAGGAGATCCTGAGATTATGTG 17750

RESULT 8
HSJ906P16      86719 bp DNA linear PRI 10-MAR-2001
LOCUS
DEFINITION Human DNA sequence from clone RP5-906P16 on chromosome
20q13.12-13.2 Contains ESTs, STSS and GSSs, complete sequence.
ACCESSION AL079339
VERSION AL079339.11 GI:6010222
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 86719)
AUTHORS Sehra,H.
TITLE Direct Submission
JOURNAL Submitted (07-MAR-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
requests: clonerequest@sanger.ac.uk
COMMENT On Oct 4, 1999 this sequence version replaced gi:5918464.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em., EMBL; Sw., SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was finished as follows unless otherwise noted: all regions were
either double-stranded or sequenced with an alternate chemistry or
covered by high quality data (i.e., phred quality >= 30); an
attempt was made to resolve all sequencing problems, such as
compressions and repeats; all regions were covered by at least one
plasmid subclone or more than one M13 subclone; and the assembly
was confirmed by restriction digest. This sequence was generated
from part of bacterial clone contigs of human chromosome 20,
constructed by the Sanger Centre Chromosome 20 Mapping Group.
Further information can be found at
http://www.sanger.ac.uk/HGP/Chr20
RP5-906P16 is from the library RPCI-5 constructed by the group of
Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pCYPAC2
IMPORTANT: This sequence is not the entire insert of clone
RP5-906P16 it may be shorter because we sequence overlapping
sections only once, except for a 100 base overlap.
The true left end of clone RP5-906P16 is at 1 in this sequence. The
true left end of clone RP5-1009H6 is at 86620 in this sequence. The
true right end of clone RP5-1009H6 is at 86620 in this sequence.
The true right end of clone RP5-1106N18 is at 58370 in this sequence.
Location/Qualifiers
1..86719
source

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/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosomes="20"
/map="q13.12-13.2"
/clone="RP5-906P16"
/clone_lib="RPCI-5"
repeat_region
415..881
/note="L1ME repeat: matches 5318..5783 of consensus"
repeat_region
1170..1730
/note="L1MB5 repeat: matches 5588..6184 of consensus"
repeat_region
1885..2135
/note="AluJo repeat: matches 2..278 of consensus"
repeat_region
2136..2187
/note="26 copies 2 mer ac 84% conserved"
repeat_region
2602..2763
/note="MIR repeat: matches 107..262 of consensus"
repeat_region
2779..3035
/note="L2 repeat: matches 2245..2481 of consensus"
repeat_region
3626..3716
/note="MIR repeat: matches 28..126 of consensus"
repeat_region
3891..4163
/note="MLT1B repeat: matches 1..246 of consensus"
repeat_region
4164..4488
/note="AluX repeat: matches 5..311 of consensus"
repeat_region
4489..4624
/note="MLT1B repeat: matches 246..388 of consensus"
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complement(4971..5437)
/note="match: GSS: Em:AQ415388"
misc_feature
complement(5094..5437)
/note="match: GSS: Em:AQ402893"
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/note="match: GSS: Em:AQ379417"
misc_feature
complement(5136..5605)
/note="match: GSS: Em:AQ135679"
repeat_region
5275..5352
/note="MIR repeat: matches 65..142 of consensus"
repeat_region
5438..5600
/note="MIR repeat: matches 91..262 of consensus"
repeat_region
5627..5921
/note="L2 repeat: matches 2398..2750 of consensus"
repeat_region
5922..6035
/note="MER58A repeat: matches 1..8 of consensus"
repeat_region
6036..6348
/note="AluSg repeat: matches 1..309 of consensus"
repeat_region
6349..6481
/note="MER58A repeat: matches 8..223 of consensus"
repeat_region
6482..6522
/note="L2 repeat: matches 2362..2398 of consensus"
misc_feature
complement(6523..6665)
/note="match: GSS: Em:B44279"
repeat_region
6666..7162
/note="LTR19A repeat: matches 1..486 of consensus"
repeat_region
7627..7717
/note="MIR repeat: matches 97..196 of consensus"
repeat_region
7720..7935
/note="MER58 repeat: matches 1..2462 of consensus"
repeat_region
8075..8323
/note="MIR repeat: matches 2..260 of consensus"
repeat_region
8456..8766
/note="L1PA4 repeat: matches 5836..6146 of consensus"
repeat_region
8767..8814
/note="24 copies 2 mer aa 83% conserved"
repeat_region
9106..9235
/note="MIR repeat: matches 9..138 of consensus"
repeat_region
9856..9940
/note="L2 repeat: matches 2624..2709 of consensus"
repeat_region
9954..10663
/note="LTR28 repeat: matches 300..1688 of consensus"
repeat_region
10634..10704
/note="LTR1 repeat: matches 119..190 of consensus"
repeat_region
10729..10893
/note="MER61B repeat: matches 1..106 of consensus"
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repeat_region
10963..11243
/note="L2 repeat: matches 2217..2496 of consensus"
repeat_region
11391..11522
/note="MER4D repeat: matches 824..972 of consensus"
misc_feature
complement(11523..12124)
/note="match: GSS: Em:AQ265342"
repeat_region
11524..12038
/note="MER4A2 repeat: matches 16..502 of consensus"
misc_feature
complement(12039..12124)
/note="match: GSS: Em:AQ279178"
repeat_region
12125..12530
/note="MER4D repeat: matches 388..793 of consensus"
repeat_region
12531..12760
/note="MER4D repeat: matches 1..234 of consensus"
repeat_region
13775..14083
/note="AluSg repeat: matches 1..309 of consensus"
repeat_region
15616..15816
/note="MER58A repeat: matches 1..203 of consensus"
repeat_region
16071..16175
/note="LTR33 repeat: matches 100..214 of consensus"
repeat_region
16411..16604
/note="L1ME repeat: matches 5595..5786 of consensus"
repeat_region
16645..17075
/note="MER83 repeat: matches 1..448 of consensus"
repeat_region
17376..17461
/note="L1ME3A repeat: matches 6048..6145 of consensus"
repeat_region
17462..17768
/note="AluJo repeat: matches 1..302 of consensus"
repeat_region
17769..17785
/note="L1ME3A repeat: matches 6145..6159 of consensus"
repeat_region
17791..17932
/note="MSTA repeat: matches 172..217 of consensus"
repeat_region
17968..18576
/note="MER4B repeat: matches 1..569 of consensus"
repeat_region
18582..18629
/note="MSTB repeat: matches 172..217 of consensus"
repeat_region
18623..18688
/note="MSTB repeat: matches 360..426 of consensus"
repeat_region
18709..19070
/note="MER61B repeat: matches 3..396 of consensus"
repeat_region
19147..19374
/note="MER46A repeat: matches 1..236 of consensus"
repeat_region
19567..19655
/note="MIR repeat: matches 78..169 of consensus"
misc_feature
complement(19656..20051)
/note="match: GSS: Em:AQ587104"
misc_feature
complement(19725..20051)
/note="match: GSS: Em:AQ123463"
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20052..20254
/note="MER20 repeat: matches 4..210 of consensus"
misc_feature
complement(20255..20679)
/note="match: GSS: Em:AQ382186"
misc_feature
complement(20301..20679)
/note="match: GSS: Em:AQ709243"
misc_feature
complement(20341..20679)
/note="match: GSS: Em:AQ121387"
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20680..20850
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misc_feature
21477..21878
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repeat_region
22066..22198
/note="MIR repeat: matches 12..156 of consensus"
repeat_region
22274..22572
/note="AluSg repeat: matches 1..292 of consensus"
repeat_region
22588..22841
/note="L2 repeat: matches 2357..2608 of consensus"
repeat_region
23153..23449
/note="MLT1E repeat: matches 245..566 of consensus"
repeat_region
23485..23536
/note="26 copies 2 mer ag 75% conserved"
repeat_region
23580..23723
/note="MLT1E repeat: matches 1..126 of consensus"
repeat_region
24119..24179
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Query Match 88.7%; Score 20.4; DB 9; Length 86719;
 Best Local Similarity 95.5%; Pred. No. 9.7;
 Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 repeat_region /note="L2 repeat: matches 2688. .2749 of consensus"
 24380. .24429 /note="25 copies 2 mer to 72% conserved"

QY 1 CAGGAGATCTGAGATTATG 22
 |||||
 Db 11441 CAGGAGATCTGAGATTATG 11462

RESULT 9
 AC079174
 LOCUS
 DEFINITION Homo sapiens 12 BAC RP11-651L5 (Roswell Park Cancer Institute Human BAC library) complete sequence.
 AC079174
 AC079174.20 GI:14277179
 HTG.
 SOURCE Homo sapiens (human)
 ORGANISM
 REFERENCE
 AUTHORS
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
 Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C., Alsbrooks,S.L., Amarantunga,H.C., Are,J.R., Banks,T., Barbara,J., Benton,J., Bimaga,K., Blankenburg,K., Bonnin,D., Bouck,J., Bowie,S., Brieva,M., Brown,M., Brown,N.P., Bryant,N.P., Buhay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C., Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Z., Chiu,D., Chowdhry,I., Christopoulos,C., Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinik,H.H., Douthwaite,K.J., Draper,H., Dugan-Rocha,S., Durbin,K.J., Earnhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Emerling,S., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T., Garza,N., Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S., Hamilton,K., Han,J., Harris,C., Harris,K., Hart,M., Haviak,P., Hawes,A., Hernandez,J., Hernandez,O., Howard,A., Hogue,M., Holloway,C., Hollins,B., Homsai,F., Howard,S., Huber,J., Hulyk,S., Hume,J., Ioshikhes,I., Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudah,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvah,J., Kovar,C., Kratovic,J., Kuresni,A., Landry,N., Leal,B., Lee,E., Lewis,L.C., Lewis,L., Li,J., Li,Z., Lichtarge,O., Lieu,C., Liu,J., Liu,W., Loulseged,H., Lozada,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R., Ma,J., Maheshwari,M., Mapua,P., Marondel,I., Martin,R., Martindale,A., Martinez,E., Massey,E., Mawhney,E., McLeod,M.P., Meador,M., Mei,G., Merscher,S., Metzker,M., Miller,A., Miner,G., Minner,Z., Mitchell,T., Mohabbat,K., Montgomery,K.T., Morgan,M., Morris,S., Moser,M., Neal,D., Nelson,D., Newton,S., Newton,N., Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokkwo,S., Oguh,M., Okuwon,G., Oragunye,N., Oviedo,R., Pace,A., Payton,B., Peery,J., Perez,L., Peters,L., Pickens,R., Primus,E., Pu,L.L., Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojibokan,I., Rolfe,M., Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shim,C., Shoohtari,N., Sison,I., Sodergren,E., Sonaike,T., Sparks,A., Stanley,H., Stone,H., Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Tamerisa,K., Tang,H., Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S., Umani,K., Vasquez,L., Vera,V., Villalon,D., Vinson,R., Wall,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S., Williams,G., Williamson,A., Wlecyk,R., Wooden,S., Worley,K., Wu,C., Wu,X., Wu,Y.F., Zhou,J., Zorrilla,S., Kucherlapati,R., Weinstein,G. and Gibbs,R.
 Direct Submission
 Unpublished
 2 (bases 1 to 87817)
 Worley,K.C.
 Submitted (23-AUG-2000) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 3 (bases 1 to 87817)
 Worley,K.C.
 Direct Submission
 Submitted (01-JUN-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 4 (bases 1 to 87817)
 Worley,K.C.
 Direct Submission
 Submitted (05-JUN-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 5 (bases 1 to 87817)
 Worley,K.C.
 Direct Submission
 Submitted (07-MAR-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 6 (bases 1 to 87817)
 Worley,K.C.
 Direct Submission
 Submitted (31-MAY-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 7 (bases 1 to 87817)
 Worley,K.C.
 Direct Submission
 Submitted (24-JUN-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 8 (bases 1 to 87817)
 Worley,K.C.
 Direct Submission
 Submitted (20-AUG-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 On Jun 1, 2001 this sequence version replaced gi:14150295.
 INFORMATION: http://www.hgsc.bcm.tmc.edu/ or email gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STSS are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found

at URL:
http://gc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html.

QUALSTAT-REPORT-----

----- Summary Statistics -----
Contig length: 189340
Phrap values in estimate: 187457
Average error rate (BCM-Phrap estimate): 3.38563e-05
Fraction of Phrap values less than 40 : 0.00584134
Number of consensus changing edits: 44
Number of N's in consensus : 0

----- Consensus changing edits -----
Position Original+Context Edited+Context
576 tcttagtttt(n)agaattaagg tcttagtttt(a)agaattaagg
7641 cctctctctt(n)gcccaataca cctctctctt(g)gcccaataca
7719 ctggtgctagt(n)ctgtacattt ctggtgctagt(c)ctgtacattt
7817 tgaagaaca(n)gactgagctg tgaagaaca(a)gactgagctg
19224 gaggacttgc(n)gccgngcng gaggacttgc(c)gccgngcng
19230 ttgngccgc(n)gngcngcng ttgngccgc(c)gngcngcng
19233 cngccgcgc(n)gccgngcng cngccgcgc(c)gccgngcng
19240 ngngccgc(n)gccgngcng ngngccgc(c)gccgngcng
22772 cactaggcca(n)gctggtggc cactaggcca(g)gctggtggc
23955 ggaatgagta(n)gtgacatttt ggaatgagta(t)gtgacatttt
46008 atggtcattga(n)atggtcactg atggtcattga(g)atggtcactg
54974 ctaactacat(n)cacagctcca ctaactacat(c)cacagctcca
57253 attatatat(n)tnnnnnntta attatatat(a)ttttattta
57255 ttatatant(n)nnnnnttat ttatatant(t)ttttattta
57256 tatatatnn(n)nnnttat tatatatnn(t)ttttattta
57257 atatatnn(n)nnnttat atatatnn(t)ttttattta
57258 tatatnnnn(n)nttatata tatatnnnn(a)ttttattta
57260 tatnnnnnn(n)nttatata tatnnnnnn(t)ttttattta
57310 tatatttta(n)atattatata tatatttta(t)atattatata
58321 ataatatttg(n)atttttagta ataatatttg(t)atttttagta
58360 gttgggcagg(n)tggtctcgaa gttgggcagg(c)tggtctcgaa
58403 acctcagct(n)ccaaagtgt acctcagct(c)ccaaagtgt
58412 tnccaagt(n)tggnntaca tnccaagt(c)tggnntaca
58417 aagtgntggg(n)ttacaggcat aagtgntggg(a)ttacaggcat
58457 aactatttgg(n)attccnaga aactatttgg(t)attccnaga
58464 tggnaattca(n)agacattatt tggnaattca(a)agacattatt
58477 acattattgt(n)tgactctgt acattattgt(t)tgactctgt
75443 taaggaggga(n)acgtagaaat taaggaggga(g)acgtagaaat
79437 aaaaagca(n)ntatagaaa aaaaagca(a)ntatagaaa
79438 aaaaagcan(n)tnatagaaa aaaaagcan(a)tnatagaaa
79440 aaaaagcant(n)atagaaa aaaaagcant(a)atagaaa
83879 ttttttttt(n)ctgntntaa ttttttttt(t)ctgntntaa
83880 ttttttttt(n)ctgntntaa ttttttttt(t)ctgntntaa
83885 ttttnctgt(n)ntaaatagt ttttnctgt(t)ntaaatagt
83886 ttttnctgt(n)ntaaatagt ttttnctgt(t)ntaaatagt
87370 tttctggagt(n)actggtcac tttctggagt(c)actggtcac
105935 aaggaattat(n)tcagaacctc aaggaattat(t)tcagaacctc
132128 tggtagact(n)tgactctgc tggtagact(c)tgactctgc
142249 tggttatttg(n)ttttttttt tggttatttg(t)ttttttttt
144054 cccaggttca(n)gngattcacc cccaggttca(a)gngattcacc
144056 cagggtcaag(n)gattcaccca cagggtcaag(t)gattcaccca
144466 tatctcatg(n)catattacat tatctcatg(t)catattacat
180955 acgattgtaa(a)aggggttttt acgattgtaa(g)aggggttttt

----- Distribution of Quality < 40 Bases -----
1000
900
800
700
Query Match 88.7%; Score 20.4; DB 9; Length 87817;
Best Local Similarity 95.5%; Pred. No. 9.7;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
DB 38408 CAGGAGATCCTGAGAAATATGTG 38429

RESULT 10
HSA132411 88665 bp DNA linear HTG 22-JUN-2001
LOCUS Homo sapiens chromosome 3p21.3, *** SEQUENCING IN PROGRESS ***, 6
DEFINITION unordered pieces.
ACCESSION AJ132411
VERSION AJ132411.1 GI:6599072
KEYWORDS HTG; HTGS_PHASE1.
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Yang, Y., Kiss, H., Kost-Alimova, M., Kedra, D., Fransson, I.,
Serousi, E., Li, J., Szeles, A., Khododnyuk, I., Imreh, M.P., Fodor, K.,
Hadlaczky, G., Klein, G., Dumaneki, J.P. and Imreh, S.
TITLE A 1-Mb PAC contig spanning the common eliminated region 1 (CER1) in
microcell hybrid-derived SCID tumors
JOURNAL Genomics 62 (2), 147-155 (1999)
MEDLINE 20079145
PUBMED 10610706
REFERENCE 2
AUTHORS Kiss, H., Kedra, D., Yang, Y., Kost-Alimova, M., Kiss, C., O'Brien, K.P.,
Fransson, I., Klein, G., Imreh, S. and Dumaneki, J.P.
TITLE A novel gene containing LIM domains (LIMD1) is located within the
common eliminated region 1 (C1CER1) in 3p21.3
JOURNAL Hum. Genet. 105 (6), 552-559 (1999)
MEDLINE 20112414
PUBMED 10647888
REFERENCE 3 (bases 1 to 88665)
AUTHORS Kedra, D.
TITLE Direct Submission
JOURNAL Submitted (15-JAN-1999) Kedra D., Dept. of Molecular Medicine,
Karolinska Hospital, Building L-8.00, Center of Molecular Medicine
(CMM), S-171 76, SWEDEN
COMMENT This sequence is unfinished. It consists of six contigs. The
contigs are
separated by gaps of unknown length. Gaps in the sequence are
represented
by a stretch of 200 NNNN. . . . All contigs in this submission are in
their
correct order and orientation. 1 22904 contig 1 23105 28324
contig 2 28525 36847 contig 3 37048 40370 contig 4 40571 54712 contig 5
54913
88665 contig 6.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 22904: contig of 22904 bp in length
* 22905 23104: gap of 200 bp
* 23105 28324: contig of 5220 bp in length
* 28325 28524: gap of 200 bp
* 28525 36847: contig of 8323 bp in length
* 36848 37047: gap of 200 bp
* 37048 40370: contig of 3323 bp in length
* 40371 40570: gap of 200 bp
* 40571 54712: contig of 14142 bp in length
* 54713 54912: gap of 200 bp
* 54913 88665: contig of 33753 bp in length.
FEATURES
Location/Qualifiers
1..88665
source

ORIGIN

Query Match 88.7%; Score 20.4; DB 2; Length 88665;
Best Local Similarity 95.5%; Pred. No. 9.7;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
Db 54284 CAGGAGATCCTGAGATTATGTG 54305

RESULT 11
AL157403

LOCUS
DEFINITION Homo sapiens chromosome 1 clone RP5-1158110 map p22.3-31.2, 13 HTG 10-JUL-2001
unordered pieces.

ACCESSION AL157403
VERSION AL157403.3 GI:9796550
KEYWORDS HTG; HTGS_PHASE1; HTGS_CANCELLED.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
1 McIay, K.
Direct Submission
TITLE Submitted (08-JUL-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
JOURNAL requests: clonerequests@sanger.ac.uk
COMMENT On Aug 12, 2000 this sequence version replaced gi:9212375.
----- Genome Center
Center: Sanger Centre
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk
----- Project Information
Center project name: dj1158110
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Sequencing vector: plasmid; L08752; 100% of reads
Chemistry: Dye-terminator ABI; 0% of reads
Chemistry: Dye-terminator Big Dye; 99% of reads
Consensus quality: 10565 bases at least Q40
Consensus quality: 107614 bases at least Q30
Consensus quality: 108618 bases at least Q20
Insert size: 109826; sum-of-contigs
Quality coverage: 3.50x in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 13 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence.
* as soon as it is available and the accession number will
* be preserved.

1 7499: contig of 7499 bp in length
* 7500: gap of 100 bp
* 10440: contig of 2841 bp in length
* 10441: 10540: gap of 100 bp
* 10541: 17763: contig of 7223 bp in length
* 17764: 17863: gap of 100 bp
* 17864: 33875: contig of 16012 bp in length
* 33876: 33975: gap of 100 bp
* 33976: 43661: contig of 9686 bp in length
* 43662: 43761: gap of 100 bp
* 43762: 52351: contig of 8590 bp in length
* 52352: 52451: gap of 100 bp
* 52452: 57434: contig of 4983 bp in length

57435: gap of 100 bp
64699: contig of 7165 bp in length
64700: gap of 100 bp
67779: contig of 2980 bp in length
67780: gap of 100 bp
75753: contig of 7874 bp in length
75754: gap of 100 bp
84741: contig of 8888 bp in length
84841: gap of 100 bp
101025: contig of 16184 bp in length
101026: gap of 100 bp
111026: contig of 9901 bp in length.

FEATURES
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/db_xref="taxon:9606"
/chromosomes="1"
/map="p22.3-31.2"
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vector_side:left"
7600..10440
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10541..17763
/note="assembly_fragment:00136
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/note="assembly_fragment:01078
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43762..52351
/note="assembly_fragment:00050"
52452..57434
/note="assembly_fragment:00066"
57535..64699
/note="assembly_fragment:00599"
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/note="assembly_fragment:01030"
67880..75753
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84842..101025
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fragment_chain:3
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vector_side:right"

ORIGIN

Query Match 88.7%; Score 20.4; DB 2; Length 111026;
Best Local Similarity 95.5%; Pred. No. 9.7;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
Db 55354 CAGGAGATCCTGAGATTATGTG 55375

RESULT 12
HS74M1/c
LOCUS HS74M1 113956 bp DNA linear PRI 27-SEP-2000

```

DEFINITION      Human DNA sequence from clone RP1-74M1 on chromosome 1p34.3-36.13
                  Contains the EPHB2 gene for a protein tyrosine kinase. Contains
                  ESTs, GSSs, STSs and a CpG island, complete sequence.
ACCESSION       AL035704
VERSION         GI:6165330
KEYWORDS        HTG; CpG island; EPHB2; tyrosine kinase.
SOURCE          Homo sapiens (human)
ORGANISM        Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 113956)
Direct Submission
Submitted (27-SEP-2000) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
requests: clonerequest@sanger.ac.uk
On Oct 31, 1999 this sequence version replaced gi:6065897.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence has been finished according to sequence map criteria
as follows. An attempt is made to resolve all sequencing problems,
such as compressions and repeats, but not necessarily within known
annotated human repeat sequence elements (e.g. Alu). Where the
sequence is ambiguous, there is an annotation using the 'unsure'
feature key.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em: EMBL; Sw: SWISSPROT; Tr: TrEMBL; Wp: WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep
This sequence was generated from part of bacterial clone configs of human
chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr1
RP1-74M1 is from the library RPCI-1 constructed at the Roswell Park
Cancer Institute by the group of Pieter de Jong. For further
details see http://bacpac.med.buffalo.edu/
VECTOR: pCYPAC2
This sequence is the entire insert of clone RP1-74M1.

FEATURES             Location/Qualifiers
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                     /organism="Homo sapiens"
                     /mol_type="genomic DNA"
                     /db_xref="taxon:9606"
                     /chromosomes="1"
                     /map="p14.3-36.13"
                     /clone="RP1-74M1"
                     /clone_lib="RPCI-1"
                     30..107
repeat_region       /note="MIR repeat: matches 39. .130 of consensus"
repeat_region       527..624
                     /note="MIR repeat: matches 47. .151 of consensus"
repeat_region       804..861
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repeat_region       880..1007
                     /note="AluJb repeat: matches 4. .125 of consensus"
repeat_region       1008..1294
                     /note="AluX repeat: matches 3. .282 of consensus"
repeat_region       1295..1439
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repeat_region       1496..1779
                     /note="AluSg repeat: matches 1. .286 of consensus"
repeat_region       1915..1956
                     /note="L1MB5 repeat: matches 6129. .6172 of consensus"
gene               2246..35193
                     /gene="EPHB2"
mRNA               join(<2246..2370,12771..12933,15359..15467,16298..16365,
25874..25996,26597..26844,27840..28055,28909..29058,
30269..30462,32331..32486,33349..33451,33545..35193)
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/product="dJ74M1.1.1 (tyrosine kinase isoform 1)"
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Em:X91737 Em:D31661 Em:L36643 Em:L43620 Em:M62325
Em:L43621 Em:L41939 Em:Z19110 Em:L25890 Em:X76011
Em:D37827 Em:M59814 Em:E09831 Em:Z19061 Em:U11493
Em:U23783 Em:Z49086 Em:U06834 Em:X65138 Em:S57168
Em:U07695 Em:Z19059 Em:AJ005029 Em:X13411 Em:D38174
Em:Z49085 Em:D14717 Em:AJ236868 Em:AB025542 Em:M59371
Em:U07634 Em:D14717 Em:AJ005026 Em:X76010 Em:X78339
Em:Z19058 Em:AB025543 Em:U03910 Em:X76012 Em:U89379
Em:M83941 Em:AB025543 Em:M68514 Em:AJ236867 Em:X60380
Em:AB040892 Em:X59290 Em:X15345 Em:L14782 Em:X52822
Em:L14823 Em:L14445 Em:AF041811 Em:L14446 Em:L14447
Em:AF176552 Em:AF216772 Em:AF216773 Em:AF245114
Em:AF216799 Em:AF237766
match: ESTs: Em:AI940384 Em:AW062316 Em:AW351853 Em:M85491
Em:AI940301 Em:AW374328 Em:AA341538 Em:T04965 Em:AW374262
Em:AA298037 Em:AA569391 Em:AI742214 Em:AW662549
Em:AA552367 Em:AI038197 Em:W96473 Em:W96506 Em:AI809403
Em:AA387292 Em:AA505599 Em:AA573757 Em:AI216739
Em:AI739658 Em:W73189 Em:AI445236 Em:AA622626 Em:AI268257
Em:AW162602 Em:AA505740 Em:AA552072 Em:W72792 Em:AI345860
Em:W76439 Em:AI702993 Em:AI476586 Em:AW365103 Em:AA570007
Em:AW168439 Em:AI344182 Em:AI911816 Em:AI345870
Em:X374261 Em:AA161293 Em:AA524449 Em:AW582672
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30269..30462,32331..32486,33349..33451,33545..35193)
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/product="dJ74M1.1.2 (tyrosine kinase isoform 2)"
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Em:AF037331 Em:M59814 Em:E09831 Em:Z19061 Em:U23783
Em:Z49086 Em:U11493 Em:E11714 Em:Z19059 Em:X65138
Em:U06834 Em:X13411 Em:D38174 Em:AJ005029 Em:Z49085
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Em:AA505559 Em:AA515992 Em:AI742214"
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30269..30462,32331..32486,33349..33451,33545..35193)
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Sw:P54763 Sw:P29323 Sw:P54762 Tr:Q91736 Sw:P09759
Tr:Q07494 Tr:Q07498 Sw:P54754 Tr:Q91735 Sw:P54753
Tr:Q95143 Sw:Q07497 Sw:P21709 Sw:Q05688 Tr:Q9Y130
Tr:Q9Y1Y3 Tr:Q9PWR5 Sw:P08069 Tr:Q9V4E5 Sw:P54761
Sw:P29317 Tr:Q9XZL6 Tr:Q9UBV3 Tr:Q9VE52 Tr:Q9UBV5

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CDS


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Sw:Q03145 Tr:Q91694 Tr:Q91776 Sw:P07948 Tr:Q9PVU9
Tr:Q91845 Tr:P89459 Sw:Q03484 Sw:P08125 Tr:O08644
Tr:O73876 Sw:P29319 Sw:Q02466 Sw:P10936 Sw:P25911
Sw:O15197 Tr:Q64736 Tr:Q952C0 Tr:Q90284 Tr:Q13332
Tr:Q63130 Tr:Q62662 Tr:Q61431 Tr:Q9VUC8 Tr:Q63132
Tr:Q60705 Sw:P41239 Tr:Q90823 Sw:Q60750 Sw:Q03127
Tr:Q9QVY9 Tr:Q9XHL4 Tr:Q93596 Tr:Q93597 Sw:P24062
Tr:Q90601 Tr:Q9XHQ4 Tr:Q26299 Tr:Q9U5A8 Tr:Q99082
Sw:P08922 Tr:Q73875 Tr:Q91433 Tr:Q42362 Tr:Q79112
Tr:Q15850 Sw:Q07407 Sw:Q04736 Tr:P79726 Tr:Q73878
Tr:Q73798 Tr:Q9U5Y2 Tr:Q15220 Tr:Q9XE13 Tr:Q13064
Sw:Q02080 Tr:Q13147 Tr:Q9VD94 Sw:Q03137 Tr:Q9PVU0
Sw:P03324 Tr:Q9PFW1 Tr:Q9PFW2 Tr:O81485 Sw:Q62413
Tr:Q28639 Sw:P32577 Sw:P00529 Tr:Q9VX92 Tr:Q9UHQ6
Sw:P54758 Sw:P00527 Tr:Q24145 Tr:Q43516 Sw:Q93105
Tr:Q64483 Sw:P54760 Tr:Q07701 Tr:Q62214 Sw:P07949
Sw:P14616 Sw:P54764 Tr:Q90478 Tr:P78440 Tr:Q63184
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Sw:P07332 Sw:P18106 Tr:Q92FS8 Tr:Q9U8W2 Tr:P93604
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Sw:P15208 Sw:P15127 Tr:Q70438 Tr:Q54951 Tr:Q9VQY1
Sw:Q07014 Sw:P41241 Sw:Q25197 Sw:P41240 Tr:Q9VHE8
Tr:Q9V9V5 Sw:P42689 Tr:Q02742 Tr:Q49202 Sw:Q29000
Sw:P42687 Tr:Q43718 Sw:Q64632 Tr:Q27576 Tr:Q54967
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join(<2248. .2370,12771. .12933,15359. .15467,16298. .16365,
25874. .25996,26597. .26844,27840. .28055,28909. .29058,
30269. .30462,32331. .32486,33349. .33457)
/gene="EPHB2"
/note="continues in dJ61A9 (AL035703), gene dJ61A9.1
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Query Match 88.7%; Score 20.4; DB 9; Length 113956;
Best Local Similarity 95.5%; Pred. No. 9.7;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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Qy 1 CAGGAGATCCTGAGATTATGTG 22
Db 94966 CAGGAGATCCTGAGATTATGTG 94945
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RESULT 13
AC106732/c AC106732 119012 bp DNA linear PRI 27-MAR-2002
LOCUS Homo sapiens chromosome 5 clone CTD-2275D24, complete sequence.
DEFINITION AC106732
ACCESSION AC106732
VERSION AC106732.2 GI:19747160
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 119012)
DOE Joint Genome Institute and Stanford Human Genome Center.
Direct Submission
Unpublished
2 (bases 1 to 119012)
DOE Joint Genome Institute.
Direct Submission
Submitted (12-JAN-2002) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
3 (bases 1 to 119012)
DOE Joint Genome Institute.
Direct Submission
Submitted (07-MAR-2002) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
4 (bases 1 to 119012)
DOE Joint Genome Institute and Stanford Human Genome Center.
Direct Submission
Submitted (27-MAR-2002) DOE Joint Genome Institute, 2800 Mitchell
Drive, Walnut Creek, CA 94598, USA
On Mar 27, 2002 this sequence version replaced gi:18139282.
Draft Sequence Produced by DOE Joint Genome Institute
www.jgi.doe.gov
Finishing Completed at Stanford Human Genome Center
www.shgc.stanford.edu
Quality: Phrap Quality >=40 99.8% of Sequence;
Estimated Total Number of Errors is 0.5.
FEATURES
source
1..119012
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTD-2275D24"
ORIGIN
Query Match 88.7%; Score 20.4; DB 9; Length 119012;
Best Local Similarity 95.5%; Pred. No. 9.7;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 CAGGAGATCCTGAGATTATGTG 22
Db 72868 CAGGAGATCCTGAGATTATGTG 72847
RESULT 14
AC002416/c AC002416 128915 bp DNA linear PRI 29-JAN-1998
LOCUS Human Chromosome X, complete sequence.
DEFINITION AC002416
ACCESSION AC002416.1 GI:2822140
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 128915)
Chen,E., Brownstein,B.H., States,D.J., Schlessinger,D. and
Mazzarella,R.
Direct Submission
Unpublished (1997)
2 (bases 1 to 128915)
Brownstein,B.H., States,D.J. and Mazzarella,R.
Direct Submission
Submitted (12-AUG-1997) Center for Genetics in Medicine, Box 8232,
Washington University School of Medicine, 4566 Scott Avenue, St.
Louis, MO 63110, USA
3 (bases 1 to 128915)
Brownstein,B.H., States,D.J. and Mazzarella,R.
Direct Submission
Submitted (29-JAN-1998) Center for Genetics in Medicine, Box 8232,
Washington University School of Medicine, 4566 Scott Avenue, St.
Louis, MO 63110, USA
```

COMMENT

On Jan 29, 1998 this sequence version replaced gi:2323255.
 Current status of this project is available at:
 'http://www.ibc.wustl.edu/cgm/seq_projects.html'
 Submitted by:
 Ellison Chen,
 Advanced Center for Genetic Technology,
 Applied Biosystems Division of Perlin Elmer Corp.,
 850 Lincoln Center Drive,
 Foster City, CA 94404 USA
 e-mail: ellison@genseq.apldbio.com

and

Buddy Brownstein,
 Center for Genetics in Medicine,
 Washington University School of Medicine, Box 8232
 4566 Scott Avenue,
 St. Louis, MO 63110, USA
 e-mail: buddy@genetics.wustl.edu

and

David J. States,
 Institute for Biomedical Computing
 Washington University in St. Louis
 700 South Euclid Ave.
 St. Louis, MO 63108 USA
 e-mail: states@ibc.wustl.edu.

FEATURES

source
 1. .128915
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"

ORIGIN

Query Match 88.7%; Score 20.4; DB 9; Length 128915;
 Best Local Similarity 95.5%; Pred. No. 9.7;
 Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCTCGAGATTATGTG 22

Db 105596 CAGGAGATCTCGAGATTATGTG 105575

RESULT 15
 AC068311/c

LOCUS AC068311 139203 bp DNA linear HTG 31-MAY-2000
 DEFINITION Homo sapiens chromosome 3 clone RP11-109P19 map 3p, WORKING DRAFT
 SEQUENCE, 47 unordered pieces.

ACCESSION AC068311

VERSION AC068311.3 GI:8121291

KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.

SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS

Cai, T., Dong, X., Gao, Q., Gu, J., Gong, J., He, F., Kang, N., Lu, T.,
 Ma, Q., Rong, L., Shen, Y., Tan, X., Wang, H., Xi, Y., Xu, Y., Yao, Z.,
 Zheng, Z., Zhu, N., Zhou, X., Zhou, Y. and Qiang, B.

Chromosome 3p genomic sequence

Unpublished

2 (bases 1 to 139203)

REFERENCE

AUTHORS

Wang, J., Hu, S., Dong, W., Wang, J., Zhang, Y., Zhang, H., Liu, B.,
 Bao, W., Sun, Y., Wu, Q., Wang, H., Yang, X., Cheng, C., Wang, Y., Niu, Y.,
 Qi, X., Li, T., Zhang, H., Liu, N., Wu, D., Yu, B., Fan, H., Liu, Y.,
 Li, G., Li, C., Bao, Q., Bao, J., Wang, X., Song, L., Zhang, L., Guo, D.,
 Huang, F., Zhang, G., Li, J., Bian, X., Zhang, M., Li, L., Feng, X., Yu, J.
 and Yang, H.

Direct Submission

JOURNAL

Submitted (02-MAY-2000) Human Genomic Center, Institute of
 Genetics, Chinese Academy of Sciences, Datun Road, Beijing, Beijing
 100101, P.R.China

COMMENT

On May 31, 2000 this sequence version replaced gi:8050886.
 -----Genome Center
 Center:Beijing Center
 Center code:Beijing
 Website:http://hgc.igtp.ac.cn
 http://www.genomics.org.cn
 Contact:hgc@igtp.ac.cn
 ----- Project Information
 Center project name:11% project
 Center clone name: RP11-109P19
 ----- Summary Statistics
 Sequencing vector: pUC18; 100% of reads
 Chemistry: Dye-terminator; ET 55% of reads
 Assembly program: Phrap; version 0.990329
 Consensus quality: 142729 bases at least Q40
 Consensus quality: 157582 bases at least Q30
 Consensus quality: 163059 bases at least Q20
 Insert size: 105278; sum-of-contigs
 Quality coverage: 7.00x in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
 * consists of 47 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

* 1 1213: contig of 1213 bp in length
 * 1214 1313: gap of unknown length
 * 1314 2442: contig of 1129 bp in length
 * 2443 2542: gap of unknown length
 * 2543 4207: contig of 1665 bp in length
 * 4208 4307: gap of unknown length
 * 4308 5648: contig of 1341 bp in length
 * 5649 5748: gap of unknown length
 * 5749 6968: contig of 1220 bp in length
 * 6969 7068: gap of unknown length
 * 7069 8123: contig of 1055 bp in length
 * 8124 8223: gap of unknown length
 * 8224 9393: contig of 1170 bp in length
 * 9394 9493: gap of unknown length
 * 9494 10687: contig of 1194 bp in length
 * 10688 10787: gap of unknown length
 * 10788 12037: contig of 1250 bp in length
 * 12038 12137: gap of unknown length
 * 12138 14344: contig of 2207 bp in length
 * 14345 14444: gap of unknown length
 * 14445 16007: contig of 1563 bp in length
 * 16008 16107: gap of unknown length
 * 16108 18196: contig of 2089 bp in length
 * 18197 18296: gap of unknown length
 * 18297 19846: contig of 1550 bp in length
 * 19847 19947: gap of unknown length
 * 19947 21757: contig of 1811 bp in length
 * 21758 21857: gap of unknown length
 * 21858 23938: contig of 2081 bp in length
 * 23939 24038: gap of unknown length
 * 24039 26931: contig of 2893 bp in length
 * 26932 27031: gap of unknown length
 * 27032 28895: contig of 1864 bp in length
 * 28896 28995: gap of unknown length
 * 28996 30602: contig of 1607 bp in length
 * 30603 30702: gap of unknown length
 * 30703 32133: contig of 1331 bp in length
 * 32034 32133: gap of unknown length
 * 32134 33637: contig of 1504 bp in length
 * 33638 33737: gap of unknown length
 * 33738 36656: contig of 2919 bp in length
 * 36657 39152: contig of 2396 bp in length
 * 39153 39252: gap of unknown length

*	3253	40461: contig of 1209 bp in length
*	40462	40561: gap of unknown length
*	40562	42590: contig of 2029 bp in length
*	42591	42690: gap of unknown length
*	42691	45244: contig of 2554 bp in length
*	45245	45344: gap of unknown length
*	45346	47973: contig of 2629 bp in length
*	47974	48073: gap of unknown length
*	48074	49724: contig of 1651 bp in length
*	49725	49824: gap of unknown length
*	49826	52090: contig of 2266 bp in length
*	52091	52190: gap of unknown length
*	52191	54464: contig of 2274 bp in length
*	54465	54564: gap of unknown length
*	54566	58061: contig of 3497 bp in length
*	58062	58161: gap of unknown length
*	58162	61037: contig of 2876 bp in length
*	61038	61137: gap of unknown length
*	61139	63773: contig of 2636 bp in length
*	63774	63873: gap of unknown length
*	63874	66477: contig of 2604 bp in length
*	66478	66577: gap of unknown length
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*	71565	71664: gap of unknown length
*	71666	75013: contig of 3349 bp in length
*	75014	75113: gap of unknown length
*	75114	79238: contig of 4125 bp in length
*	79239	79338: gap of unknown length
*	79339	82528: contig of 3190 bp in length
*	82529	82628: gap of unknown length
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*	87636	87735: gap of unknown length
*	87736	93180: contig of 5445 bp in length
*	93181	93280: gap of unknown length
*	93281	98141: contig of 4861 bp in length
*	98142	98241: gap of unknown length
*	98242	104076: contig of 5835 bp in length
*	104077	104176: gap of unknown length
*	104177	110939: contig of 6763 bp in length
*	110940	111039: gap of unknown length
*	111041	115184: contig of 4145 bp in length
*	115185	115284: gap of unknown length
*	115285	121746: contig of 6462 bp in length
*	121747	121846: gap of unknown length
*	121847	129746: contig of 7900 bp in length
*	129747	129846: gap of unknown length
*	129847	139203: contig of 9357 bp in length

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misc_feature 1314. .2442 /note="assembly_name:Contig75"  
misc_feature 2343. .4207 /notes="assembly_name:Contig84"  
misc_feature 4308. .5648 /note="assembly_name:Contig86"  
misc_feature 5749. .6968 /notes="assembly_name:Contig87"  
misc_feature 7069. .8123 /notes="assembly_name:Contig89"  
misc_feature 8224. .9393 /notes="assembly_name:Contig90"  
misc_feature 9494. .10687 /notes="assembly_name:Contig92"
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misc_feature	12138. .14344 /note="assembly_name:Contig94"
misc_feature	14445. .16007 /note="assembly_name:Contig95"
misc_feature	16108. .18196 /note="assembly_name:Contig96"
misc_feature	18297. .19846 /note="assembly_name:Contig97"
misc_feature	19947. .21757 /note="assembly_name:Contig98"
misc_feature	21858. .23938 /note="assembly_name:Contig99"
misc_feature	24039. .26931 /note="assembly_name:Contig100"
misc_feature	27032. .28895 /note="assembly_name:Contig101"
misc_feature	28996. .30602 /note="assembly_name:Contig102"
misc_feature	30703. .32033 /note="assembly_name:Contig103"
misc_feature	32134. .33637 /note="assembly_name:Contig104"
misc_feature	33738. .36656 /note="assembly_name:Contig105"
misc_feature	36757. .39152

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Query Match      88.7%; Score 20.4; DB 2; Length 139203;
Best Local Similarity 95.5%; Pred. No. 9.7;
Matches 21; Conservative 0; Mismatches 1; Indels 0;
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Search completed: August 13, 2005, 05:04:37
Job time : 738.234 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 12, 2005, 22:21:57 ; Search time 184.245 Seconds
(without alignments)
738.985 Million cell updates/sec

Title: US-10-673-854-4

Perfect score: 23

Sequence: 1 caggagatcctgagattatgttg 23

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

N Geneseq_16Dec04:*
1: Geneseqn1980s:*
2: Geneseqn1990s:*
3: Geneseqn2000s:*
4: Geneseqn2001as:*
5: Geneseqn2001bs:*
6: Geneseqn2002as:*
7: Geneseqn2002bs:*
8: Geneseqn2003as:*
9: Geneseqn2003bs:*
10: Geneseqn2003cs:*
11: Geneseqn2003ds:*
12: Geneseqn2004as:*
13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	19.8	86.1	2995	10	Adf82489 Leukaemia
2	18.8	81.7	1112	10	Adc20846 Human sec
3	18.8	81.7	1112	10	Abt16983 Human sec
4	18.8	81.7	1112	10	Abz67931 Human sec
5	18.8	81.7	1149	12	Adn05709 Antipsori
6	18.8	81.7	1160	3	Aaf22348 Human sec
7	18.8	81.7	1160	10	Adc20223 Human sec
8	18.8	81.7	1160	10	Abt16829 Human sec
9	18.8	81.7	1160	10	Abz67143 Human sec
10	18.8	81.7	1238	10	Adc20845 Human sec
11	18.8	81.7	1238	10	Abt16982 Human sec
12	18.8	81.7	1238	10	Abz67930 Human sec
13	18.8	81.7	1545	4	Aak86702 Human imm
14	18.8	81.7	1545	4	Aak86701 Human imm
15	18.8	81.7	1547	4	Aak86700 Human imm
16	18.8	81.7	2158	8	Abx34674 Human mdd
17	18.8	81.7	13919	6	Abk86218 DNA encod
18	18.8	81.7	13919	6	Abk86221 AIP-1/FLA
19	18.8	81.7	13919	6	Abk86220 AIP-1/FLA
20	18.8	81.7	21404	6	Abk86229 AIP-1/FLA

21	18.8	81.7	55795	6	ABL68242	Abt168242 Kidney ca
22	18.8	81.7	55795	6	ABL68863	Abt168863 Kidney ca
23	18.8	81.7	55795	6	ABL68484	Abt168484 Kidney ca
24	18.8	81.7	55795	6	ABN95045	Abn95045 Gene #154
25	18.8	81.7	58320	13	ABD33125	Abd33125 Human can
26	18.8	81.7	110000	10	ADG70447_0	Adg70447 Human ANG
27	18.8	81.7	110000	10	ABZ79565_0	Abz79565 CLLD8 and
28	18.8	81.7	188794	12	ADQ59476	Adq59476 Human can
29	18.8	81.7	226215	11	ACN45146	Acn45146 Human gen
30	18.2	79.1	1097	12	ADL90243	Adl90243 Human gen
31	18.2	79.1	1220	12	ADL90240	Adl90240 Human enz
32	18.2	79.1	2311	12	ADQ67262	Adq67262 Novel hum
33	18.2	79.1	2797	6	ABE57465	Abt57465 Human pro
34	18.2	79.1	3243	12	ADL90241	Adl90241 Human enz
35	18.2	79.1	3359	12	ADL90242	Adl90242 Human enz
36	18.2	79.1	3367	9	AAD57344	Aad57344 Human kin
37	18.2	79.1	138115	13	ABD33312	Abd33312 Murine ca
38	17.8	77.4	700	4	AHH92092	Aah92092 Human inf
39	17.8	77.4	42024	12	ADQ97373	Adq97373 Human can
40	17.8	77.4	110000	3	AAF22303_2	Continuation (3 of
41	17.2	74.8	135	5	ABA16804	Abat16804 Human ner
42	17.2	74.8	140	4	AAK58947	Aak58947 Human imm
43	17.2	74.8	148	12	ACH92736	Ach92736 Human gen
44	17.2	74.8	149	4	AAK58234	Aak58234 Human imm
45	17.2	74.8	203	2	AAT24266	Aat24266 Human gen

ALIGNMENTS

RESULT 1

ADf82489
ID ADF82489 standard; DNA; 2995 BP.
XX
AC ADF82489;
XX
DT 26-FEB-2004 (first entry)
XX
DE Leukaemia-related DNA sequence #3045.
XX
KW Cytostatic; Gene therapy; leukaemia; ss.
XX
OS Unidentified.
XX
PN WO2003039443-A2.
XX
PD 15-MAY-2003.
XX
PF 04-NOV-2002; 2002WO-EF012303.
XX
PR 05-NOV-2001; 2001EP-00126244.
PR 30-APR-2002; 2002EP-00009758.
XX

(DEKR-) DEUT KREBSFORSCHUNGSZENTRUM.

(UYLU-) UNIV LUDWIG MAXIMILIANS.

(HAFE/) HAERLACH T.

(SCHG/) SCHOCH C.

(KERN/) KERN W.

Haerlach T, Schoch C, Kern W, Kohlmann A, Schnittger S, Dugas M;

Bills R, Brors B, Mergenthaler S;

WPI; 2003-505037/47.

Determining the subtype of leukemia cells and whether a patient sample

contains leukemia cells or other cells, useful for treating leukemia,

comprises determining the expression profile of a group of markers in a

patient sample.

Disclosure; SEQ ID NO 3045; 2938pp; English.

The present invention relates to a method (M1) for determining the

subtype of leukaemia cells and whether a patient sample contains

CC leukaemia cells. The method comprises determining the expression profile
 CC of a group of markers in a patient sample. The method is useful for
 CC determining the presence of leukaemia cells, its types or subtypes, and
 CC for the preparation of a medicament for treating leukaemia.

XX SQ Sequence 2995 BP; 845 A; 585 C; 700 G; 865 T; 0 U; 0 Other;
 Query Match 86.1%; Score 19.8; DB 10; Length 2995;
 Best Local Similarity 91.3%; Pred. No. 19;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 23
 ID 2029 CAGGAATCTGAGACTATGTG 2051
 DB

RESULT 2
 ADC20846
 ID ADC20846 standard; DNA; 1112 BP.
 XX AC ADC20846;
 XX DT 18-DEC-2003 (first entry)
 XX DE Human secreted protein-related DNA sequence #264.
 XX KW gene therapy; human; secreted protein; haemopoietic disorder;
 KW haematological disorder; anaemia; haemophilia; inflammatory disorder;
 KW inflammatory bowel disease; Crohn's disease; neoplastic disease; cancer;
 KW leukaemia; wound healing; epithelial cell proliferation disorder;
 KW immune disorder; autoimmune disorder; asthmatic disorder;
 KW cardiovascular disorder; atherosclerosis; myocarditis;
 KW infectious disease; HIV; AIDS; endocrine disorder; diabetes;
 KW gastrointestinal disorder; duodenal ulcer; gastroenteritis; gene; ds.
 XX OS Homo sapiens.
 XX PN WO200292787-A2.
 XX PD 21-NOV-2002.
 XX PF 26-MAR-2002; 2002WO-US009257.
 XX PR 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX PA Rosen CA, Ruben SM;
 XX PI WPI; 2003-129287/12.
 XX DR
 XX PT New human secreted proteins and nucleic acid molecules, useful for
 PT preparing a diagnostic or pharmaceutical composition for diagnosing,
 PT preventing or treating hematopoietic or hematologic disorders, e.g.
 PT anemia or hemophilia.
 XX PS Disclosure; SEQ ID NO 800; 1512pp; English.
 XX CC The invention comprises the amino acid and coding sequences of human
 CC secreted proteins. The DNA and protein sequences of the invention are
 CC useful for detecting, preventing, diagnosing, prognosticating, treating
 CC or ameliorating: haematopoietic or haematological disorders (e.g. anaemia
 CC and haemophilia); inflammatory disorders (e.g. inflammatory bowel disease
 CC and Crohn's disease); neoplastic disease (e.g. cancer and leukaemia);
 CC wound healing and disorders of epithelial cell proliferation; immune
 CC disorders (e.g. autoimmune disorders and asthmatic disorders);
 CC cardiovascular disorders (e.g. atherosclerosis and myocarditis);
 CC infectious disease (e.g. HIV/AIDS); endocrine disorders (e.g. diabetes);
 CC and gastrointestinal disorders (e.g. duodenal ulcers and
 CC gastroenteritis). The present DNA sequence was used in the
 CC exemplification of the invention.

XX SQ Sequence 1112 BP; 267 A; 221 C; 178 G; 446 T; 0 U; 0 Other;
 Query Match 81.7%; Score 18.8; DB 10; Length 1112;
 Best Local Similarity 90.9%; Pred. No. 50;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
 ID 369 CAGGAGATCCTGAGAACATGTG 390
 DB

RESULT 3
 ABT16983
 ID ABT16983 standard; DNA; 1112 BP.
 XX AC ABT16983;
 XX DT 03-APR-2003 (first entry)
 XX DE Human secreted protein-related DNA sequence - SEQ ID No 337.
 XX KW Human; gene; ds; protein therapy; immediate hypersensitivity disease;
 KW allergic disorder; asthmatic disorder; gene therapy; secreted protein;
 KW hay fever; allergic conjunctivitis; allergic rhinitis;
 KW binding partner identification; chromosome identification;
 KW radiation hybrid mapping; long-range restriction mapping.
 XX OS Homo sapiens.
 XX PN WO200277188-A2.
 XX PD 03-OCT-2002.
 XX PF 26-MAR-2002; 2002WO-US009239.
 XX PR 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX PA Rosen CA, Ruben SM;
 XX PI WPI; 2003-175010/17.
 XX DR
 XX PT Use of human secreted proteins and nucleic acids for preparing a
 PT diagnostic or pharmaceutical composition for diagnosing or treating
 PT allergic or asthmatic disorders, e.g. asthma, hay fever, or allergic
 PT conjunctivitis or rhinitis.
 XX PS Disclosure; Page 769; 823pp; English.
 XX CC The invention comprises the amino acid and coding sequences of human
 CC secreted proteins. The DNA and protein sequences of the invention are
 CC useful for the diagnosis and treatment of allergic disorders, asthmatic
 CC disorders and immediate hypersensitivity diseases (e.g. hay fever,
 CC allergic conjunctivitis and allergic rhinitis). The proteins of the
 CC invention are also useful for identifying a binding partner. The nucleic
 CC acids of the invention are also useful for chromosome identification,
 CC radiation hybrid mapping or long-range restriction mapping. The present
 CC DNA sequence represents a human secreted protein-related DNA sequence

XX SQ Sequence 1112 BP; 267 A; 221 C; 178 G; 446 T; 0 U; 0 Other;
 Query Match 81.7%; Score 18.8; DB 10; Length 1112;
 Best Local Similarity 90.9%; Pred. No. 50;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
 ID 369 CAGGAGATCCTGAGAACATGTG 390
 DB

```
RESULT 4
ABZ67931
ID ABZ67931 standard; DNA; 1112 BP.
XX
XX
AC ABZ67931;
XX
XX
DT 26-MAR-2003 (first entry)
XX
XX
DE Human secreted protein encoding genomic DNA SEQ ID NO 1454.
XX
XX
XX Human; secreted protein; neutropenic; neuroprotective; cytostatic;
KW virucide; dermatological; immunosuppressive; anti-inflammatory;
KW vulnary; antibacterial; antiparkinsonian; antiscikling; antianemic;
KW antiarthritic; cancer; antirheumatic; hepatotropic; cerebroprotective;
KW antiinflammatory; anti-allergic; antidiabetic; antiulcer; anticonvulsant;
KW antifungal; antiparasitic; cardiant; immune disorder; infection; vaccine;
KW cardiovascular disorder; neurological disease; nephrotropic;
KW gene therapy; gene; ds.
XX
XX OS Homo sapiens.
XX
XX PN WO20027186-A2.
XX
XX PD 03-OCT-2002.
XX
XX PF 26-MAR-2002; 2002WO-US009188.
XX
XX PR 27-MAR-2001; 2001US-027850P.
XX
XX PR 12-SEP-2001; 2001US-00950082.
XX
XX PR 12-SEP-2001; 2001US-00950083.
XX
XX PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX PI Rosen CA, Ruben SM;
XX
XX DR WPI; 2003-040583/03.
XX
XX PT New human secreted proteins encoded by genes contained in cDNA clones
PT (e.g. HGCAC19), useful for preventing, treating or diagnosing e.g. AIDS,
PT multiple sclerosis, herpes virus, leukemia, tick-borne encephalitis or
PT West Nile fever.
XX
XX PS Disclosure; Page 2064; 2423pp; English.
XX
XX CC The invention relates to novel human genes (ABZ66891-ABZ68209) and the
XX encoded secreted proteins (ABP99470-ABP99872) useful for preventing,
XX treating or ameliorating medical conditions e.g. by protein or gene
XX therapy. The genes are isolated from a range of human tissues disclosed
XX in the specification. The nucleic acids, proteins, antibodies and
XX (ant)agonists are useful in the diagnosis, treatment and prevention of:
XX (a) cancer, e.g. breast and ovarian cancer and other cancers of the
XX adrenal gland, bone, bone marrow, breast, gastrointestinal tract, liver,
XX lung or urogenital; (b) immune disorders e.g. Addison's disease,
XX allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis,
XX diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid
XX arthritis and ulcerative colitis; (c) cardiovascular disorders such as
XX myocardial ischaemias; (d) wound healing; (e) neurological diseases e.g.
XX cerebral anoxia and epilepsy; and (f) infectious diseases such as viral,
XX bacterial, fungal and parasitic infections
XX
XX SQ Sequence 1112 BP; 267 A; 221 C; 178 G; 446 T; 0 U; 0 Other;

Query Match 81.7%; Score 18.8; DB 10; Length 1112;
Best Local Similarity 90.9%; Pred. No. 50;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 369 CAGGAGATCCTGAGAACATGTG 390

RESULT 5
ABZ67931
ID ABZ67931 standard; DNA; 1112 BP.
XX
XX
AC ABZ67931;
XX
XX
DT 26-MAR-2003 (first entry)
XX
XX
DE Human secreted protein encoding genomic DNA SEQ ID NO 1454.
XX
XX
XX Human; secreted protein; neutropenic; neuroprotective; cytostatic;
KW virucide; dermatological; immunosuppressive; anti-inflammatory;
KW vulnary; antibacterial; antiparkinsonian; antiscikling; antianemic;
KW antiarthritic; cancer; antirheumatic; hepatotropic; cerebroprotective;
KW antiinflammatory; anti-allergic; antidiabetic; antiulcer; anticonvulsant;
KW antifungal; antiparasitic; cardiant; immune disorder; infection; vaccine;
KW cardiovascular disorder; neurological disease; nephrotropic;
KW gene therapy; gene; ds.
XX
XX OS Homo sapiens.
XX
XX PN WO20027186-A2.
XX
XX PD 03-OCT-2002.
XX
XX PF 26-MAR-2002; 2002WO-US009188.
XX
XX PR 27-MAR-2001; 2001US-027850P.
XX
XX PR 12-SEP-2001; 2001US-00950082.
XX
XX PR 12-SEP-2001; 2001US-00950083.
XX
XX PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX PI Rosen CA, Ruben SM;
XX
XX DR WPI; 2003-040583/03.
XX
XX PT New human secreted proteins encoded by genes contained in cDNA clones
PT (e.g. HGCAC19), useful for preventing, treating or diagnosing e.g. AIDS,
PT multiple sclerosis, herpes virus, leukemia, tick-borne encephalitis or
PT West Nile fever.
XX
XX PS Disclosure; Page 2064; 2423pp; English.
XX
XX CC The invention relates to novel human genes (ABZ66891-ABZ68209) and the
XX encoded secreted proteins (ABP99470-ABP99872) useful for preventing,
XX treating or ameliorating medical conditions e.g. by protein or gene
XX therapy. The genes are isolated from a range of human tissues disclosed
XX in the specification. The nucleic acids, proteins, antibodies and
XX (ant)agonists are useful in the diagnosis, treatment and prevention of:
XX (a) cancer, e.g. breast and ovarian cancer and other cancers of the
XX adrenal gland, bone, bone marrow, breast, gastrointestinal tract, liver,
XX lung or urogenital; (b) immune disorders e.g. Addison's disease,
XX allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis,
XX diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid
XX arthritis and ulcerative colitis; (c) cardiovascular disorders such as
XX myocardial ischaemias; (d) wound healing; (e) neurological diseases e.g.
XX cerebral anoxia and epilepsy; and (f) infectious diseases such as viral,
XX bacterial, fungal and parasitic infections
XX
XX SQ Sequence 1112 BP; 267 A; 221 C; 178 G; 446 T; 0 U; 0 Other;

Query Match 81.7%; Score 18.8; DB 10; Length 1112;
Best Local Similarity 90.9%; Pred. No. 50;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 369 CAGGAGATCCTGAGAACATGTG 390
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```
ADN05709
ID ADN05709 standard; cDNA; 1149 BP.
XX
XX AC ADN05709;
XX
XX DT 01-JUL-2004 (first entry)
XX
XX DE Antipsoriatic cDNA sequence #1084.
XX
XX KW ds; gene; antipsoriatic; gene therapy; psoriasis; diagnosis.
XX
XX OS Homo sapiens.
XX
XX PN WO2004028479-A2.
XX
XX PD 08-APR-2004.
XX
XX PF 25-SEP-2003; 2003WO-US030907.
XX
XX PR 25-SEP-2002; 2002US-0414006P.
XX
XX (GETH ) GENENTECH INC.
XX
XX Bodary S, Clark H, Jackman J, Schoenfeld J, Williams PM, Wood WT;
XX Wu TD;
XX
XX WPI; 2004-305105/28.
XX
XX P-PSDB; ADN05710.
XX
XX New PRO nucleic acid or polypeptide, useful for preparing a
XX pharmaceutical composition for diagnosing or treating psoriasis in a
XX mammal.
XX
XX Claim 1; SEQ ID NO 2103; 3069pp; English.
XX
XX The invention relates to novel polynucleotide and polypeptides for
XX treating psoriasis or a sequence having at least 80% identity to the
XX above sequences. The nucleic acid is useful for preparing a composition
XX for diagnosing or treating psoriasis in a mammal. This sequence
XX corresponds to one of the polynucleotides of the invention.
XX
XX SQ Sequence 1149 BP; 276 A; 219 C; 184 G; 420 T; 0 U; 50 Other;

Query Match 81.7%; Score 18.8; DB 12; Length 1149;
Best Local Similarity 90.9%; Pred. No. 51;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 367 CAGGAGATCCTGAGAACATGTG 388

RESULT 6
AAF22348
ID AAF22348 standard; cDNA; 1160 BP.
XX
XX AC AAF22348;
XX
XX DT 26-MAR-2001 (first entry)
XX
XX DE Human secreted protein gene 33 SEQ ID NO:43.
XX
XX KW Human; secreted protein; diagnosis; immunosuppressive; antiarthritic;
XX antirheumatic; antiproliferative; cytostatic; cardiant; vasotropic;
XX cerebroprotective; neutropenic; neuroprotective; antibacterial; virucide;
XX fungicide; ophthalmological; vulnary; gene therapy; neoplasm;
XX autoimmune disease; rheumatoid arthritis; hyperproliferative disorder;
XX cardiovascular disorder; cardiac arrest; cerebrovascular disorder;
XX cerebral ischaemia; angiogenesis; nervous system disorder; infection;
XX Alzheimer's disease; ocular disorder; corneal infection; wound healing;
XX skin aging; food additive; preservative; ss.
XX
XX OS Homo sapiens.
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XX WO200061748-A1.
 PN 19-OCT-2000.
 XX 06-APR-2000; 2000WO-US008982.
 XX 09-APR-1999; 99US-0128696P.
 PR 14-JAN-2000; 2000US-0176069P.
 XX (HUMA-) HUMAN GENOME SCI INC.
 PA Rosen CA, Ruben SM, Komatsoulis G;
 PI WPI; 2000-638566/61.
 XX P-PSDB; AAB63081.
 DR New nucleic acid molecules encoding 48 human secreted proteins for
 PT diagnosing, preventing, treating or ameliorating medical conditions and
 PT used as food additives or preservatives.
 XX
 PS Claim 1; Page 421; 480pp; English.
 XX
 CC AAF22316 to AAF22363 encode the human secreted proteins given in AAB63049
 CC to AAB63096. AAB63097 to AAB63132 represent more human secreted proteins
 CC and polypeptides homologous to them. Human secreted proteins have
 CC activities based on the tissues and cells the genes are expressed in.
 CC Examples of activities include: immunosuppressive; antiarthritic;
 CC antirheumatic; antiproliferative; cytostatic; cardiant; vasotropic;
 CC cerebroprotective; neurotropic; neuroprotective; antibacterial; virucide;
 CC fungicide; ophthalmological; and vulnerary. The polynucleotides and
 CC proteins can be used to prevent, treat or ameliorate a medical condition
 CC in e.g. humans, mice, rabbits, goats, horses, cats, dogs, chickens or
 CC sheep. They are also used in diagnosing a pathological condition or
 CC susceptibility to a pathological condition. Disorders which are diagnosed
 CC or treated include autoimmune diseases e.g. rheumatoid arthritis,
 CC hyperproliferative disorders e.g. neoplasms of the breast or liver,
 CC cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders
 CC e.g. cerebral ischaemia, angioneurosis, nervous system disorders e.g.
 CC Alzheimer's disease, infections caused by bacteria, viruses and fungi and
 CC ocular disorders e.g. corneal infection. The polypeptides can also be
 CC used to aid wound healing and epithelial cell proliferation, to prevent
 CC skin aging due to sunburn, to maintain organs before transplantation, for
 CC supporting cell culture of primary tissues, to regenerate tissues and in
 CC chemotaxis. The polypeptides can also be used as a food additive or
 CC preservative to increase or decrease storage capabilities, fat content,
 CC lipid, protein, carbohydrate, vitamins, minerals, cofactors and other
 CC nutritional components. AAF22307 to AAF22315 and AAB63048 represent
 CC sequences used in the exemplification of the present invention
 XX
 SQ Sequence 1160 BP; 294 A; 227 C; 184 G; 453 T; 0 U; 2 Other;
 Query Match 81.7%; Score 18.8; DB 3; Length 1160;
 Best Local Similarity 90.9%; Pred. No. 51;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 CAGGAGATCCTGAGATTATGTG 22
 Db |||||
 370 CAGGAGATCCTGAGACATGTG 391
 RESULT 7
 ADC20223
 ID ADC20223 standard; DNA; 1160 BP.
 XX
 AC ADC20223;
 XX
 DT 18-DEC-2003 (first entry)
 XX Human secreted protein coding sequence #162.
 DE gene therapy; human; secreted protein; haemopoietic disorder;
 KW haematological disorder; anaemia; haemophilia; inflammatory disorder;
 KW

KW inflammatory bowel disease; Crohn's disease; neoplastic disease; cancer;
 KW leukaemia; wound healing; epithelial cell proliferation disorder;
 KW immune disorder; autoimmune disorder; asthmatic disorder;
 KW cardiovascular disorder; atherosclerosis; myocarditis;
 KW infectious disease; HIV; AIDS; endocrine disorder; diabetes;
 KW gastrointestinal disorder; duodenal ulcer; gastroenteritis; gene; ds.
 XX Homo sapiens.
 OS WO200292787-A2.
 XX 21-NOV-2002.
 XX 26-MAR-2002; 2002WO-US009257.
 XX 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX (HUMA-) HUMAN GENOME SCI INC.
 PA Rosen CA, Ruben SM;
 XX WPI; 2003-129287/12.
 DR New human secreted proteins and nucleic acid molecules, useful for
 PT preparing a diagnostic or pharmaceutical composition for diagnosing,
 PT preventing or treating hematopoietic or hematologic disorders, e.g.
 PT anemia or hemophilia.
 XX Claim 1; SEQ ID NO 172; 1512pp; English.
 PS The invention comprises the amino acid and coding sequences of human
 XX secreted proteins. The DNA and protein sequences of the invention are
 CC useful for detecting, preventing, diagnosing, prognosticating, treating
 CC or ameliorating: hematopoietic or hematological disorders (e.g. anaemia
 CC and haemophilia); inflammatory disorders (e.g. inflammatory bowel disease
 CC and Crohn's disease); neoplastic disease (e.g. cancer and leukaemia);
 CC wound healing and disorders of epithelial cell proliferation; immune
 CC disorders (e.g. autoimmune disorders and asthmatic disorders);
 CC cardiovascular disorders (e.g. atherosclerosis and myocarditis);
 CC infectious disease (e.g. HIV/AIDS); endocrine disorders (e.g. diabetes);
 CC and gastrointestinal disorders (e.g. duodenal ulcers and
 CC gastroenteritis). The present DNA sequence encodes a human secreted
 CC protein of the invention.
 XX Sequence 1160 BP; 294 A; 227 C; 184 G; 453 T; 0 U; 2 Other;
 SQ Query Match 81.7%; Score 18.8; DB 10; Length 1160;
 Best Local Similarity 90.9%; Pred. No. 51;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 CAGGAGATCCTGAGATTATGTG 22
 Db |||||
 370 CAGGAGATCCTGAGAACATGTG 391
 RESULT 8
 ABT16829
 ID ABT16829 standard; DNA; 1160 BP.
 XX
 AC ABT16829;
 XX
 DT 03-APR-2003 (first entry)
 XX Human secreted protein gene sequence - SEQ ID No 78.
 DE Human; gene; ds; protein therapy; immediate hypersensitivity disease;
 XX allergic disorder; asthmatic disorder; gene therapy; secreted protein;
 KW hay fever; allergic conjunctivitis; allergic rhinitis;
 KW binding partner identification; chromosome identification;
 KW radiation hybrid mapping; long-range restriction mapping.
 XX


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OS Homo sapiens.
PN WO200277188-A2.
XX
XX
PD 03-OCT-2002.
XX
XX
PF 26-MAR-2002; 2002WO-US009239.
XX
XX
PR 27-MAR-2001; 2001US-0278650P.
PR 12-SEP-2001; 2001US-00950082.
PR 12-SEP-2001; 2001US-00950083.
XX
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX
PI Rosen CA, Ruben SM;
XX
XX WPI; 2003-175010/17.
XX
XX Use of human secreted proteins and nucleic acids for preparing a
PT diagnostic or pharmaceutical composition for diagnosing or treating
PT allergic or asthmatic disorders, e.g. asthma, hay fever, or allergic
PT conjunctivitis or rhinitis.
XX
XX
PS Claim 7; Page 601-602; 823pp; English.
XX
XX The invention comprises the amino acid and coding sequences of human
CC secreted proteins. The DNA and protein sequences of the invention are
CC useful for the diagnosis and treatment of allergic disorders, asthmatic
CC disorders and immediate hypersensitivity diseases (e.g. hay fever,
CC allergic conjunctivitis and allergic rhinitis). The proteins of the
CC invention are also useful for identifying a binding partner. The nucleic
CC acids of the invention are also useful for chromosome identification,
CC radiation hybrid mapping or long-range restriction mapping. The present
CC DNA sequence encodes a human secreted protein of the invention
XX
XX
SQ Sequence 1160 BP; 294 A; 227 C; 184 G; 453 T; 0 U; 2 Other;

Query Match 81.7%; Score 18.8; DB 10; Length 1160;
Best Local Similarity 90.9%; Pred. No. 51;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
   |||||
DB 370 CAGGAGATCCTGAGAACATGTG 391

RESULT 9
ABZ67143
ID ABZ67143 standard; cDNA; 1160 BP.
XX
XX ABZ67143;
XX
XX 26-MAR-2003 (first entry)
XX
XX Human secreted protein encoding cDNA SEQ ID NO 263.
XX
XX Human; secreted protein; neuroprotective; cytosolic;
KW virucide; dermatological; immunosuppressive; antiinflammatory; anti-HIV;
KW vulnery; antibacterial; antiparkinsonian; antickling; antianaemic;
KW antiarthritic; cancer; antirheumatic; hepatotropic; cerebroprotective;
KW antiinflammatory; antiallergic; antidiabetic; antitumor; anticonvulsant;
KW antifungal; antiparasitic; cardiant; immune disorder; infection; vaccine;
KW cardiovascular disorder; neurological disease; nephrotropic;
KW gene therapy; gene; ds.
XX
XX Homo sapiens.
XX
XX WO200277186-A2.
XX
XX 03-OCT-2002.
XX
XX 26-MAR-2002; 2002WO-US009188.
XX
XX
PF 26-MAR-2002; 2002WO-US009188.
XX
XX

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PR 27-MAR-2001; 2001US-0278650P.
PR 12-SEP-2001; 2001US-00950082.
PR 12-SEP-2001; 2001US-00950083.
XX
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX
PI Rosen CA, Ruben SM;
XX
XX WPI; 2003-040583/03.
XX
XX P-PSDB; ABP99722.
XX
XX New human secreted proteins encoded by genes contained in cDNA clones
PT (e.g. HGCAC19), useful for preventing, treating or diagnosing e.g. AIDS,
PT multiple sclerosis, herpes virus, leukemia, tick-borne encephalitis or
PT West Nile fever.
XX
XX
XX Claim 7; Page 1308; 2423pp; English.
XX
XX The invention relates to novel human genes (ABZ66891-ABZ68209) and the
CC encoded secreted proteins (ABP99470-ABP99872) useful for preventing,
CC treating or ameliorating medical conditions e.g. by protein or gene
CC therapy. The genes are isolated from a range of human tissues disclosed
CC in the specification. The nucleic acids, proteins, antibodies and
CC (ant)agonists are useful in the diagnosis, treatment and prevention of:
CC (a) cancer, e.g. breast and ovarian cancer and other cancers of the
CC adrenal gland, bone, bone marrow, breast, gastrointestinal tract, liver,
CC lung or urogenital; (b) immune disorders e.g. Addison's disease,
CC allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis,
CC diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid
CC arthritis and ulcerative colitis; (c) cardiovascular disorders such as
CC myocardial ischaemias; (d) wound healing; (e) neurological diseases e.g.
CC cerebral anoxia and epilepsy; and (f) infectious diseases such as viral,
CC bacterial, fungal and parasitic infections
XX
XX
SQ Sequence 1160 BP; 294 A; 227 C; 184 G; 453 T; 0 U; 2 Other;

Query Match 81.7%; Score 18.8; DB 10; Length 1160;
Best Local Similarity 90.9%; Pred. No. 51;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
   |||||
DB 370 CAGGAGATCCTGAGAACATGTG 391

RESULT 10
ADC20845
ID ADC20845 standard; DNA; 1238 BP.
XX
XX ADC20845;
XX
XX 18-DEC-2003 (first entry)
XX
XX Human secreted protein-related DNA sequence #263.
XX
XX gene therapy; human; secreted protein; haemopoietic disorder;
KW haematological disorder; anaemia; haemophilia; inflammatory disorder;
KW inflammatory bowel disease; Crohn's disease; neoplastic disease; cancer;
KW leukaemia; wound healing; epithelial cell proliferation disorder;
KW immune disorder; autoimmune disorder; asthmatic disorder;
KW cardiovascular disorder; atherosclerosis; myocarditis;
KW infectious disease; HIV; AIDS; endocrine disorder; diabetes;
KW gastrointestinal disorder; duodenal ulcer; gastroenteritis; gene; ds.
XX
XX Homo sapiens.
XX
XX WO200292787-A2.
XX
XX 21-NOV-2002.
XX
XX 26-MAR-2002; 2002WO-US009257.
XX
XX 27-MAR-2001; 2001US-0278650P.

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XX The invention relates to novel human genes (ABZ66891-ABZ68209) and the
CC encoded secreted proteins (ABP9470-ABP99872) useful for preventing,
CC treating or ameliorating medical conditions e.g. by protein or gene
CC therapy. The genes are isolated from a range of human tissues disclosed
CC in the specification. The nucleic acids, proteins, antibodies and
CC (ant)agonists are useful in the diagnosis, treatment and prevention of:
CC (a) cancer, e.g. breast and ovarian cancer and other cancers of the
CC adrenal gland, bone, bone marrow, breast, gastrointestinal tract, liver,
CC lung or urogenital; (b) immune disorders e.g. Addison's disease,
CC allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis,
CC diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid
CC arthritis and ulcerative colitis; (c) cardiovascular disorders such as
CC myocardial ischaemias; (d) wound healing; (e) neurological diseases e.g.
CC cerebral anoxia and epilepsy; and (f) infectious diseases such as viral,
CC bacterial, fungal and parasitic infections
XX
SQ Sequence 1238 BP; 322 A; 231 C; 208 G; 477 T; 0 U; 0 Other;

Query Match 81.7%; Score 18.8; DB 10; Length 1238;

Best Local Similarity 90.9%; Pred. No. 51;

Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CAGGAGATCCTGAGATTATG 22

Db 369 CAGGAGATCCTGAGACATG 390

RESULT 13

AAK86702/C

ID AAK86702 standard; DNA; 1545 BP.

XX AAK86702;

AC AAK86702;

XX 07-NOV-2001 (first entry)

DT 07-NOV-2001 (first entry)

XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:41514.

DE Human; immune/haematopoietic; immune/haematopoietic antigen; cancer;

XX Human; immune/haematopoietic; immune/haematopoietic antigen; cancer;

KW cytostatic; gene therapy; vaccine; metastasis; ds.

KW cytostatic; gene therapy; vaccine; metastasis; ds.

XX Homo sapiens.

OS Homo sapiens.

XX WO200157182-A2.

PN WO200157182-A2.

XX 09-AUG-2001.

PD 09-AUG-2001.

XX 17-JAN-2001; 2001WO-US0001354.

PF 17-JAN-2001; 2001WO-US0001354.

XX 31-JAN-2000; 2000US-0179065P.

PR 04-FEB-2000; 2000US-0180628P.

PR 24-FEB-2000; 2000US-0184664P.

PR 02-MAR-2000; 2000US-0186350P.

PR 16-MAR-2000; 2000US-0189874P.

PR 17-MAR-2000; 2000US-0190076P.

PR 18-APR-2000; 2000US-0198123P.

PR 19-MAY-2000; 2000US-0205515P.

PR 07-JUN-2000; 2000US-0209467P.

PR 28-JUN-2000; 2000US-0214886P.

PR 30-JUN-2000; 2000US-0215135P.

PR 07-JUL-2000; 2000US-0216647P.

PR 07-JUL-2000; 2000US-0216880P.

PR 11-JUL-2000; 2000US-0217487P.

PR 11-JUL-2000; 2000US-0217496P.

PR 14-JUL-2000; 2000US-0218290P.

PR 26-JUL-2000; 2000US-0220963P.

PR 14-AUG-2000; 2000US-0220964P.

PR 14-AUG-2000; 2000US-0224518P.

PR 14-AUG-2000; 2000US-0224519P.

PR 14-AUG-2000; 2000US-0225213P.

PR 14-AUG-2000; 2000US-0225214P.

PR 14-AUG-2000; 2000US-0225266P.

PR 14-AUG-2000; 2000US-0225267P.

PR 14-AUG-2000; 2000US-0225268P.

PR 14-AUG-2000; 2000US-0225270P.

PR 14-AUG-2000; 2000US-0225447P.

PR 14-AUG-2000; 2000US-0225757P.

PR 14-AUG-2000; 2000US-0225758P.

PR 14-AUG-2000; 2000US-0225759P.

PR 18-AUG-2000; 2000US-0226279P.

PR 22-AUG-2000; 2000US-0226681P.

PR 22-AUG-2000; 2000US-0226868P.

PR 22-AUG-2000; 2000US-0227182P.

PR 23-AUG-2000; 2000US-0227009P.

PR 30-AUG-2000; 2000US-0228924P.

PR 01-SEP-2000; 2000US-0229287P.

PR 01-SEP-2000; 2000US-0229343P.

PR 01-SEP-2000; 2000US-0229344P.

PR 01-SEP-2000; 2000US-0229345P.

PR 05-SEP-2000; 2000US-0229509P.

PR 05-SEP-2000; 2000US-0229513P.

PR 06-SEP-2000; 2000US-0230437P.

PR 06-SEP-2000; 2000US-0230438P.

PR 08-SEP-2000; 2000US-0231242P.

PR 08-SEP-2000; 2000US-0231243P.

PR 08-SEP-2000; 2000US-0231244P.

PR 08-SEP-2000; 2000US-0231413P.

PR 08-SEP-2000; 2000US-0231414P.

PR 08-SEP-2000; 2000US-0232080P.

PR 08-SEP-2000; 2000US-0232081P.

PR 12-SEP-2000; 2000US-0231968P.

PR 14-SEP-2000; 2000US-0232397P.

PR 14-SEP-2000; 2000US-0232398P.

PR 14-SEP-2000; 2000US-0232399P.

PR 14-SEP-2000; 2000US-0232400P.

PR 14-SEP-2000; 2000US-0232401P.

PR 14-SEP-2000; 2000US-0233063P.

PR 14-SEP-2000; 2000US-0233064P.

PR 14-SEP-2000; 2000US-0233065P.

PR 21-SEP-2000; 2000US-0234223P.

PR 21-SEP-2000; 2000US-0234274P.

PR 25-SEP-2000; 2000US-0234997P.

PR 25-SEP-2000; 2000US-0234998P.

PR 26-SEP-2000; 2000US-0235484P.

PR 27-SEP-2000; 2000US-0235834P.

PR 27-SEP-2000; 2000US-0235836P.

PR 29-SEP-2000; 2000US-0236327P.

PR 29-SEP-2000; 2000US-0236367P.

PR 29-SEP-2000; 2000US-0236368P.

PR 29-SEP-2000; 2000US-0236369P.

PR 29-SEP-2000; 2000US-0236370P.

PR 02-OCT-2000; 2000US-0236802P.

PR 02-OCT-2000; 2000US-0237037P.

PR 02-OCT-2000; 2000US-0237038P.

PR 02-OCT-2000; 2000US-0237039P.

PR 02-OCT-2000; 2000US-0237040P.

PR 13-OCT-2000; 2000US-0239935P.

PR 13-OCT-2000; 2000US-0239937P.

PR 20-OCT-2000; 2000US-0240960P.

PR 20-OCT-2000; 2000US-0241221P.

PR 20-OCT-2000; 2000US-0241785P.

PR 20-OCT-2000; 2000US-0241786P.

PR 20-OCT-2000; 2000US-0241787P.

PR 20-OCT-2000; 2000US-0241808P.

PR 20-OCT-2000; 2000US-0241809P.

PR 20-OCT-2000; 2000US-0241826P.

PR 01-NOV-2000; 2000US-0244617P.

PR 08-NOV-2000; 2000US-0246474P.

PR 08-NOV-2000; 2000US-0246475P.

PR 08-NOV-2000; 2000US-0246476P.

PR 08-NOV-2000; 2000US-0246477P.

PR 08-NOV-2000; 2000US-0246478P.

PR 08-NOV-2000; 2000US-0246523P.

PR 08-NOV-2000; 2000US-0246524P.

PR 08-NOV-2000; 2000US-0246525P.

PR 08-NOV-2000; 2000US-0246526P.

PR 08-NOV-2000; 2000US-0246527P.
PR 08-NOV-2000; 2000US-0246528P.
PR 08-NOV-2000; 2000US-0246532P.
PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.
PR 08-NOV-2000; 2000US-0246611P.
PR 08-NOV-2000; 2000US-0246613P.
PR 17-NOV-2000; 2000US-0249207P.
PR 17-NOV-2000; 2000US-0249208P.
PR 17-NOV-2000; 2000US-0249209P.
PR 17-NOV-2000; 2000US-0249210P.
PR 17-NOV-2000; 2000US-0249211P.
PR 17-NOV-2000; 2000US-0249212P.
PR 17-NOV-2000; 2000US-0249213P.
PR 17-NOV-2000; 2000US-0249214P.
PR 17-NOV-2000; 2000US-0249215P.
PR 17-NOV-2000; 2000US-0249216P.
PR 17-NOV-2000; 2000US-0249217P.
PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249244P.
PR 17-NOV-2000; 2000US-0249245P.
PR 17-NOV-2000; 2000US-0249264P.
PR 17-NOV-2000; 2000US-0249265P.
PR 17-NOV-2000; 2000US-0249297P.
PR 17-NOV-2000; 2000US-0249299P.
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PR 06-DEC-2000; 2000US-0251479P.
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PR 08-DEC-2000; 2000US-0251989P.
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PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-0259678P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Barash SC, Ruben SM;
XX WPI; 2001-483426/52.
XX
XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
XX useful for preventing, diagnosing and/or treating cancers and metastasis.
XX
XX Disclosure; SEQ ID NO 41514; 3071pp + Sequence Listing; English.
XX
XX AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)
XX amino acid sequences given in AAK82170 to AAK91921. (I) have cytostatic
XX activity, and can be used in gene therapy and vaccine production. (I)
XX proteins and polynucleotides may be used in the prevention, diagnosis and
XX treatment of diseases associated with inappropriate (I) expression. For
XX example, they may be used to treat disorders associated with decreased
XX expression by rectifying mutations or deletions in a patient's genome
XX that affect the activity of (I) by expressing inactive proteins or to
XX supplement the patients own production of (I). Additionally, (I)
XX polynucleotides may be used to produce the secreted (I), by inserting the
XX nucleic acids into a host cell and culturing the cell to express the
XX protein. (I) proteins and polynucleotides may be used to prevent,
XX diagnose and treat immune/haematopoietic-related diseases, especially
XX cancers and cancer metastases of haematopoietic-derived cells. AAK64703
XX to AAK87694 represent human immune/haematopoietic antigen genomic
XX sequences from the present invention. AAK54942 to AAK54950 and AAK82169
XX represent sequences used in the exemplification of the present invention
XX
XX SQ Sequence 1545 BP; 575 A; 204 C; 311 G; 455 T; 0 U; 0 Other;

Query Match 81.7%; Score 18.8; DB 4; Length 1545;
Best Local Similarity 90.9%; Pred. No. 53;

Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 CAGGAGATCCTGAGATTATGTG 22
Db 762 CAGAAGGTCCTGAGATTATGTG 741
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ID AAK86701 standard; DNA; 1545 BP.
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XX AAK86701;
XX
XX 07-NOV-2001 (first entry)
XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:41513.
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
XX cytostatic; gene therapy; vaccine; metastasis; ds.
XX Homo sapiens.
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XX WO200157182-A2.
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XX 09-AUG-2001.
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XX 17-JAN-2001; 2001WO-US001354.
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PR 08-DEC-2000; 2000US-0251990P.
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(HUNA-) HUMAN GENOME SCI INC.
PA XX
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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Title: US-10-673-854-4

Perfect score: 23

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Searched: 1202784 seqs, 818138359 residues

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Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 3	18.8	81.7	601	4	US-09-949-016-88904
C 4	18.8	81.7	601	4	US-09-949-016-160361
C 5	18.8	81.7	16505	4	US-09-949-016-119362
C 6	18.8	81.7	16506	4	US-09-949-016-119366
C 7	18.8	81.7	25202	4	US-09-949-016-13151
C 8	18.8	81.7	30053	4	US-09-949-016-16231
C 9	18.8	81.7	47375	4	US-09-949-016-15420
C 10	18.8	81.7	58108	4	US-09-949-016-13383
C 11	18.8	81.7	62386	4	US-09-949-016-12823
C 12	18.8	81.7	62386	4	US-09-949-016-12823
C 13	18.8	81.7	93510	4	US-09-949-016-15095
C 14	18.8	81.7	112623	4	US-09-949-016-14374
C 15	18.8	81.7	162465	4	US-09-949-016-14264
C 16	18.8	81.7	234894	4	US-09-949-016-16420
C 17	18.2	79.1	57761	4	US-09-949-016-13429
C 18	17.8	77.4	601	4	US-09-949-016-134977
C 19	17.8	77.4	24954	4	US-09-949-016-13475
C 20	17.8	77.4	78157	4	US-09-949-016-16466
C 21	17.8	77.4	78157	4	US-09-949-016-16467
C 22	17.8	77.4	256176	4	US-09-949-016-12822
C 23	17.8	77.4	256176	4	US-09-949-016-15524
C 24	17.8	77.4	636591	4	US-09-949-016-11808
C 25	17.8	77.4	636591	4	US-09-949-016-11388
C 26	17.2	74.8	601	4	US-09-949-016-50649
C 27	17.2	74.8	601	4	US-09-949-016-50650

C 28	17.2	74.8	601	4	US-09-949-016-53774	Sequence 53774, A
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C 30	17.2	74.8	601	4	US-09-949-016-64795	Sequence 64795, A
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C 39	17.2	74.8	601	4	US-09-949-016-172198	Sequence 172198, A
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C 42	17.2	74.8	601	4	US-09-949-016-199070	Sequence 199070, A
C 43	17.2	74.8	601	4	US-09-949-016-204673	Sequence 204673, A
C 44	17.2	74.8	780	4	US-09-573-080A-178	Sequence 178, App
C 45	17.2	74.8	1017	4	US-09-573-080A-177	Sequence 177, App

ALIGNMENTS

RESULT 1
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; Sequence 56405, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 56405
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; ORGANISM: Human
US-09-949-016-56405

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Best Local Similarity 90.9%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
DB 520 CAGGAGATCCTGAGAACATGTG 499

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; Patent No. 6812339
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; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498

; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 56406
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-56406

Query Match 81.7%; Score 18.8; DB 4; Length 601;
Best Local Similarity 90.9%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
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DB 528 CAGGAGATCCTGAGACATGTG 507

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; Sequence 88904, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
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; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
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; TYPE: DNA
; ORGANISM: Human
US-09-949-016-160361

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; Patent No. 6812339
; GENERAL INFORMATION:
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; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
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; TYPE: DNA
; ORGANISM: Human
US-09-949-016-11962

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; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13836
; LENGTH: 16506
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13836

Query Match 81.7%; Score 18.8; DB 4; Length 16506;

Best Local Similarity 90.9%; Pred. No. 19;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 878 CAGGAGATCCTGAGAACATGTG 899

RESULT 7

US-09-949-016-13151
; Sequence 13151, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13151
; LENGTH: 25202
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13151

Query Match 81.7%; Score 18.8; DB 4; Length 25202;
Best Local Similarity 90.9%; Pred. No. 20;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 5221 CAGGAGATCCTGAGAACATGTG 5242

RESULT 8

US-09-949-016-16231
; Sequence 16231, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16231
; LENGTH: 30053
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-16231

Query Match 81.7%; Score 18.8; DB 4; Length 30053;
Best Local Similarity 90.9%; Pred. No. 20;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||

Db 20662 CAGGAGATCCTGAGAACATGTG 20683

RESULT 9

US-09-949-016-15420
; Sequence 15420, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15420
; LENGTH: 47375
; TYPE: DNA
; ORGANISM: Human
; NAME/KEY: misc feature
; LOCATION: (1)...(47375)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-15420

Query Match 81.7%; Score 18.8; DB 4; Length 47375;
Best Local Similarity 90.9%; Pred. No. 22;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 18878 CAGGAGATCCTGAGAACATGTG 18899

RESULT 10

US-09-949-016-13383
; Sequence 13383, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13383
; LENGTH: 58108
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13383

Query Match 81.7%; Score 18.8; DB 4; Length 58108;
Best Local Similarity 90.9%; Pred. No. 22;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 33903 CAGGAGATCCTGAGAACATGTG 33924

```
RESULT 11
US-09-949-016-12823
; Sequence 12823, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12823
; LENGTH: 62386
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(62386)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-12823

Query Match      81.7%; Score 18.8; DB 4; Length 62386;
Best Local Similarity 90.9%; Pred. No. 23;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 CAGGAGATCCTGAGATTATGTG 22
Db      34283 CAGGAGATCCTGAGACATGTG 34304

RESULT 12
US-09-949-016-12823/c
; Sequence 12823, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12823
; LENGTH: 62386
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(62386)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-12823

Query Match      81.7%; Score 18.8; DB 4; Length 62386;
Best Local Similarity 90.9%; Pred. No. 23;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 CAGGAGATCCTGAGATTATGTG 22
Db      34283 CAGGAGATCCTGAGACATGTG 34304

RESULT 13
US-09-949-016-15095
; Sequence 15095, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15095
; LENGTH: 93510
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(93510)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-15095

Query Match      81.7%; Score 18.8; DB 4; Length 93510;
Best Local Similarity 90.9%; Pred. No. 24;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 CAGGAGATCCTGAGATTATGTG 22
Db      42219 CAGGAGATCCTGAGAACATGTG 42240

RESULT 14
US-09-949-016-14374/c
; Sequence 14374, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14374
; LENGTH: 112623
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(112623)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14374

Query Match      81.7%; Score 18.8; DB 4; Length 112623;
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Best Local Similarity 90.9%; Pred. No. 25;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
| | | | | | | | | | | | | | | | | | | | | |
Db 75109 CAGGAGATCCTGAGACATGTG 75088

RESULT 15
US-09-949-016-14264/c
; Sequence 14264, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14264
; LENGTH: 162465
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-14264

Query Match 81.7%; Score 18.8; DB 4; Length 162465;
Best Local Similarity 90.9%; Pred. No. 26;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
| | | | | | | | | | | | | | | | | | | | | |
Db 20162 CAGGAGATCCTGAGACATGTG 20141

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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(without alignments)
592.293 Million cell updates/sec

Title: US-10-673-854-4
Perfect score: 23
Sequence: 1 caggagatcctgagattatgtgg 23

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST.*
1: gb_est1:*
2: gb_est2:*
3: gb_hic:*
4: gb_est3:*
5: gb_est4:*
6: gb_est5:*
7: gb_est6:*
8: gb_gss1:*
9: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20.4	88.7	459	AI024054	AI024054 ov72b12.s
2	20.4	88.7	605	AA719016	AA719016 ab46b02.s
3	19.8	86.1	521	BE754118	BE754118 207531.MA
4	19.4	84.3	456	AQ136698	AQ136698 HS_2195.A
5	19	82.6	417	CE041276	CE041276 tigr-gss-
6	18.8	81.7	221	AA831411	AA831411 oc60h12.s
7	18.8	81.7	257	BF806428	BF806428 RC2-CI008
8	18.8	81.7	281	AQ069577	AQ069577 HS_2267.B
9	18.8	81.7	333	BY653205	BY653205 BY653205
10	18.8	81.7	337	AQ472951	AQ472951 CITBI-E1-
11	18.8	81.7	378	AA608986	AA608986 af05d07.s
12	18.8	81.7	386	AQ279565	AQ279565 CITBI-E1-
13	18.8	81.7	401	AA569281	AA569281 nm31e01.s
14	18.8	81.7	414	AL703856	AL703856 DREZp686N
15	18.8	81.7	441	AQ022522	AQ022522 HS_2180.A
16	18.8	81.7	449	AA701122	AA701122 zg56b06.s
17	18.8	81.7	467	AQ172559	AQ172559 HS_3197.A
18	18.8	81.7	470	EX101541	EX101541 BX101541
19	18.8	81.7	475	AQ471928	AQ471928 CITBI-E1-
20	18.8	81.7	482	B94003	B94003 CIT-HSP-217
21	18.8	81.7	487	AQ428760	AQ428760 CITBI-E1-
22	18.8	81.7	488	AA708132	AA708132 zg06e08.s
23	18.8	81.7	503	AQ734095	AQ734095 HS_3109.B
24	18.8	81.7	512	AQ204899	AQ204899 HS_3226_B

C	25	18.8	81.7	523	8	AQ544890	AQ544890 CITBI-E1-
C	26	18.8	81.7	532	8	AQ684672	AQ684672 HS_5481.B
C	27	18.8	81.7	564	8	AQ149724	AQ149724 HS_3178.A
28	29	18.8	81.7	564	8	AQ316239	AQ316239 RPII11-10
29	28	18.8	81.7	565	8	AQ791207	AQ791207 HS_4555.B
30	30	18.8	81.7	573	8	AQ504120	AQ504120 RPII11-2
31	31	18.8	81.7	592	2	AW964983	AW964983 EST377056
C	32	18.8	81.7	594	8	AQ478792	AQ478792 RPII11-2
C	33	18.8	81.7	608	1	AI887916	AI887916 tk14d01.X
C	34	18.8	81.7	614	4	BI061657	BI061657 IL3-UT011
C	35	18.8	81.7	628	8	AQ485321	AQ485321 RPII11-2
C	36	18.8	81.7	629	8	AQ550526	AQ550526 RPII11-4
C	37	18.8	81.7	630	4	BI061656	BI061656 IL3-UT011
C	38	18.8	81.7	637	1	AV684105	AV684105 AV684105
C	39	18.8	81.7	638	9	AG160747	AG160747 Pan trogl
40	41	18.8	81.7	641	8	AQ742110	AQ742110 HS_5566.B
41	40	18.8	81.7	658	9	AG160363	AG160363 Pan trogl
42	42	18.8	81.7	659	4	BI062008	BI062008 IL3-UT011
43	43	18.8	81.7	663	9	CE227365	CE227365 tigr-gss-
44	44	18.8	81.7	692	5	BM990062	BM990062 UI-H-DIO-
45	45	18.8	81.7	694	7	CF993883	CF993883 AGENCOURT

ALIGNMENTS

RESULT 1
LOCUS AI024054 459 bp mRNA linear EST 27-AUG-1998
DEFINITION ov72b12.s1 Soares_testis_NHT Homo sapiens cDNA clone IMAGE:1642847
3', mRNA sequence.
ACCESSION AI024054
VERSION AI024054.1 GI:3239098
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 459)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabbs.r@mail.nih.gov
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LINL at: www.bio.llnl.gov/bbrp/image/image.html
Insert length: 1046 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 449.

FEATURES

Location/Qualifiers
1..459
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1642847"
/sex="male"
/lab_host="DH10B"
/clone_lib="Soares_testis_NHT"
/note="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was prepared from mRNA obtained from Clontech Laboratories, Inc., and primed with a Not I - oligo(dT) primer [5']
TGTTACCAATCTGAGTGGAGCGCGCCCAATTTTTTTTTTTT 3'.
Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library

went through one round of normalization to Cot5, and was constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN

Query Match 88.7%; Score 20.4; DB 1; Length 459;
Best Local Similarity 95.5%; Pred. No. 81;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||

Db 15 CAGGAGATCCTGAGATTATGTG 36
|||||

RESULT 2

AA719016 605 bp mRNA linear EST 12-JAN-1999
LOCUS ah46b02.s1 Soares_testis_NHT Homo sapiens cDNA clone 1292523 3',
DEFINITION mRNA sequence.

ACCESSION AA719016
VERSION AA719016.1 GI:2732115
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 605)
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
JOURNAL Contact: Robert Strausberg, Ph.D.
COMMENT Email: cgabbs-r@mail.nih.gov
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone Distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www.biol.llnl.gov/bbrp/image/image.html
Insert Length: 1362 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 481.

FEATURES

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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="1292523"
/sex="male"
/lab_host="DH10B"
/clone_lib="Soares_testis_NHT"
/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was prepared from mRNA obtained from Clontech Laboratories, Inc., and primed with a Not I - oligo(dT) primer [5']
TGTTACCATCTGAGTGGAGCGCGCCCAATTTTTTTTTTTT 3']
Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization to Cot5, and was constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN

Query Match 88.7%; Score 20.4; DB 1; Length 605;
Best Local Similarity 95.5%; Pred. No. 83;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||

Db 16 CAGGAGATCCTGAGATTATGTG 37
|||||

RESULT 3

BE754118/c
LOCUS BE754118
DEFINITION 207531 MARC 2BOV Bos taurus cDNA 5', mRNA sequence.
ACCESSION BE754118
VERSION BE754118.1 GI:10168110
KEYWORDS EST.
SOURCE Bos taurus (cow)
ORGANISM Bos taurus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovinae; Bos.
1 (bases 1 to 521)
Smith, T.P.L., Grosse, W.M., Freking, B.A., Roberts, A.J., Stone, R.T.,
Casas, E., Wray, J.E., White, J., Cho, J., Fahrenkrug, S.C.,
Bennett, G.L., Heaton, M.P., Laegreid, W.W., Rohrer, G.A.,
Chitko-McKown, C.G., Pettea, G., Holt, I., Karamycheva, S., Liang, F.,
Quackenbush, J., and Keefe, J.W.
Sequence evaluation of four pooled-tissue normalized bovine cDNA
libraries and construction of a gene index for cattle
Genome Res. 11 (4), 626-630 (2001)

REFERENCE

AUTHORS Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred
v0.980904.e. Vector identified by cross_match with the -minscore 18
and -minmatch 12 options.
PCR Primers
FORWARD: AGGAACACAGCTATGACCAT
BACKWARD: GTTTCCTCAGTCACGACG
Plate: 51 row: E column: 11
Seq primer: ATTGAGTGACACTATAG.
Location/Qualifiers
1..521
/organism="Bos taurus"
/mol_type="mRNA"
/db_xref="taxon:9913"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="MARC 2BOV"
/note="Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI;
Library made from pooled tissue from testis, thymus,
semitendinosus muscle, longissimus muscle, pancreas,
adrenal, and endometrium."

JOURNAL

MEDLINE

PUBMED

COMMENT

FEATURES

source

ORIGIN

Query Match 86.1%; Score 19.8; DB 2; Length 521;
Best Local Similarity 91.3%; Pred. No. 1.6e+02;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 23
|||||

Db 404 CAGGAGATCCTGAGATTCTGTGG 382
|||||

RESULT 4

AQ136698/c
LOCUS AQ136698
DEFINITION HS_2195_A2_F08_MF CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2195 Col=16 Row=K, genomic survey sequence.
ACCESSION AQ136698
VERSION AQ136698.1 GI:3523764
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 456)
 AUTHORS Mahairas,G.C., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.
 TITLE Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome
 JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
 MEDLINE 99380589
 PUBMED 1049764
 COMMENT Contact: Mahairas GG, Wallace JC, Hood L
 High Throughput Sequencing Center
 University of Washington
 401 Queen Anne Avenue North, Seattle, WA 98109, USA
 Tel: (206) 616-3618
 Fax: (206) 616-3887
 Email: jwallace@u.washington.edu
 Sequence Tagged Connector
 Plats: 2195 row: K column: 16
 Class: BAC ends
 High quality sequence stop: 456.

FEATURES
 source
 1..456
 Location/Qualifiers
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /clone="Plate=2195 Col=16 Row=K"
 /sex="male"
 /clone_lib="CIT Approved Human Genomic Sperm Library D"
 /note="Organ: sperm; Vector: pBelobAC11; BAC Clones in E-Coli DH10B"

ORIGIN

Query Match 84.3%; Score 19.4; DB 8; Length 456;
 Best Local Similarity 90.3%; Pred. No. 2.4e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22

Db 456 CAGGAGATCCTGAGATCATGTG 435

RESULT 5
 CE041276 417 bp DNA linear GSS 24-SEP-2003
 LOCUS tigr-gss-dog-17000350045460 Dog Library Canis familiaris genomic,
 DEFINITION genomic survey sequence.
 ACCESSION CE041276
 VERSION CE041276.1 GI:35073799
 KEYWORDS GSS.

SOURCE

ORGANISM Canis familiaris (dog)
 Canis familiaris
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.

REFERENCE 1 (bases 1 to 417)
 AUTHORS Kirkness,E.F., Bafna,V., Halpern,A.L., Levy,S., Remington,K., Rusch,D.B., Delcher,A.L., Pop,M., Wang,W., Fraser,C.M. and Venter,J.C.
 TITLE The dog genome: survey sequencing and comparative analysis
 JOURNAL Science 301 (5641), 1898-1903 (2003)
 MEDLINE 22875432
 PUBMED 14512627
 COMMENT Contact: Kirkness EF
 The Institute for Genomic Research
 Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
 Rockville, MD 20850, USA
 Tel: 301-838-0200
 Fax: 301-838-0208
 Email: ekirknes@tigr.org
 Class: shotgun.

FEATURES

source
 1..417
 Location/Qualifiers
 /organism="Canis familiaris"
 /mol_type="genomic DNA"

ORIGIN

Query Match 82.6%; Score 19; DB 9; Length 417;
 Best Local Similarity 100.0%; Pred. No. 3.7e+02;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 AGGAGATCCTGAGATTATG 20

Db 329 AGGAGATCCTGAGATTATG 347

RESULT 6

AA831411 221 bp mRNA linear EST 07-APR-1998
 LOCUS oc60h12.s1 NCI CGAP GCB1 Homo sapiens cDNA clone IMAGE:1354151 3'
 DEFINITION similar to contains MER4.t3 MER4 repetitive element ;, mRNA sequence.
 ACCESSION AA831411
 VERSION AA831411.1 GI:2904510
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
 JOURNAL Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman, Ph.D., Gerald Marti, M.D.
 CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.

CDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
 www.bio.llnl.gov/bbrp/image/image.html
 Insert Length: 445 Std Error: 0.00
 Seq primer: -40ml3 fwd. ET from Amersham
 High quality sequence stop: 190.

FEATURES

source

1..221
 Location/Qualifiers
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:1354151"
 /tissue_type="germinal center B cell"
 /lab_host="DH10B"
 /clone_lib="NCI CGAP GCB1"
 /note="Vector: p773D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was prepared from human tonsillar cells enriched for germinal center B cells by flow sorting (CD20+, IgD-), provided by Dr. Louis M. Staudt (NCI), Dr. David Allman (NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was primed with a Not I - oligo(dT) primer
 [5'-TGTACCAATCTGAAGTGGAGCGCCGCCATCTTTT-3'
]. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified p773 vector. Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN

Query Match 81.7%; Score 18.8; DB 1; Length 221;

Best Local Similarity 90.9%; Pred. No. 4.3e+02; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 62 CAGGAGATCCTGAGATGTG 83

RESULT 7
BF806428 257 bp mRNA linear EST 12-JAN-2001
LOCUS RC2-CI0088-081100-011-f10 CI0088 Homo sapiens cDNA, mRNA sequence.
ACCESSION BF806428
VERSION BF806428.1 GI:12135417
KEYWORDS EST.

SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 257)

AUTHORS Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R., Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F., Goldman, G.H., Carvalho, A.F., Matsukuma, A., Baia, G.S., Simpson, D.H., Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and Simpson, A.J.
COMMENT Shotgun sequencing of the human transcriptome with ORF expressed sequence tags

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

MEDLINE 20202663
PubMed 10737800
CONTACT: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001

Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=RC2&t2=RC2-CI0088-081100-011-f10&t3=2000-11-08&t4=1)
Seq primer: puc 18 forward
High quality sequence start: 25
High quality sequence stop: 257.

FEATURES
source Location/Qualifiers

1..257
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="CI0088"
/note="Organ: colon ins; Vector: puc18; Site 1: SmaI;
Site 2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the puc 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN
Query Match 81.7%; Score 18.8; DB 2; Length 257;
Best Local Similarity 90.9%; Pred. No. 4.4e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db 233 CAGGAGATCCTGAGATCTGTG 254

RESULT 8
AQ069577/c

LOCUS AQ069577 281 bp DNA linear GSS 04-AUG-1998
DEFINITION HS_2267_B2_G01_MF_CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2267 Col=2 Row=N, genomic survey sequence.

ACCESSION AQ069577
VERSION AQ069577.1 GI:3384776
KEYWORDS GSS.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 281)

AUTHORS Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T., Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and Hood, L.

TITLE Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

MEDLINE 99380589

PubMed 10449764

COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Sequence Tagged Connector

Plate: 2267 row: N column: 2

Class: BAC ends

High quality sequence stop: 281.

FEATURES
source Location/Qualifiers

1..281

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/clone_lib="Plate=2267 Col=2 Row=N"

/sex="male"

/clone_libs="CIT Approved Human Genomic Sperm Library D"

/note="Organ: sperm; Vector: pBelOBAC11; BAC Clones in E-Coli DH10B"

ORIGIN

Query Match 81.7%; Score 18.8; DB 8; Length 281;
Best Local Similarity 90.9%; Pred. No. 4.4e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||

Db 194 CAGGAGATCCTGAGACATGTG 173
|||||

RESULT 9

LOCUS BY653205 333 bp mRNA linear EST 16-DEC-2002
DEFINITION BY653205 RIKEN full-length enriched, visual cortex Mus musculus cDNA clone K330308P06 3', mRNA sequence.

ACCESSION BY653205

VERSION BY653205.1 GI:27013497

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 333)

AUTHORS Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S., Nikaido, I., Oeato, N., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C., Gojobori, T., Baldarelli, R., Hill, D.P., Bult, C., Hume, D.A., Quackenbush, J., Schriml, L.M., Kanapin, A., Matsuda, H., Batalov, S., Beisel, K.W., Blake, J.A., Bradt, D., Bruscia, V., Chothia, C., Corbani, L.E., Cousins, S., Dalla, E., Dragani, T.A., Fletcher, C.F., Forrest, A., Frazer, K.S., Gaasterland, T.,

Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I.J., Jarvis, E.D., Kanai, A., Kawai, H., Kawasawa, Y., Kedzierski, R.M., King, B.L., Konagaya, A., Kurochkin, I.V., Lee, Y., Lenhard, B., Lyons, P.A., Maglott, D.R., Maltais, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W.J., Perte, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D., Ramachandran, S., Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring, B.Z., Ringwald, M., Sandelin, A., Schneider, C., Semple, C.A., Setou, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y., Yang, L., Yuan, Z., Zavalon, M., Zhu, Y., Zimmer, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K., Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shingawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E.S., Rogers, J., Birney, E. and Hayashizaki, Y.

TITLE
Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs

JOURNAL
Nature 420, 563-573 (2002)

MEDLINE
22354683

PUBMED
12466851

COMMENT
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@sc.riken.jp, URL: http://genome.gsc.riken.jp/
Aizawa, K., Akimura, T., Aizawa, T., Carninci, P., Fukuda, S., Hirozane, T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Konno, H., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Waki, K., Watahiki, A., Muramatsu, M. and Hayashizaki, Y. Direct Submission

Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)
RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in Riken Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.
Tissues were provided by Michela Fagiolini and Takao K. Hensch (Laboratory for Neuronal Circuit Development Brain Science Institute RIKEN 2-1 Hiroosawa, Wako-shi, Saitama 351-0198 Japan) whose assistance we gratefully acknowledge.
Please visit our web site (http://genome.gsc.riken.go.jp) for further details.

FEATURES
source

Location/Qualifiers
1. .333
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="K530308P06"
/tissue_type="visual cortex"
/clone_lib="RIKEN full-length enriched, visual cortex"

ORIGIN

Query Match 81.7%; Score 18.8; DB 6; Length 333;
Best Local Similarity 90.9%; Pred. No. 4.5e+02;

Matches
20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY
1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db
104 CAGGAGATCCTGAGATTATGTG 125
|||||

RESULT 10
AQ472951
LOCUS
CITBI-El-2589A16.1 TR CITBI-El Homo sapiens genomic clone 2589A16,
genomic survey sequence.
AQ472951
DEFINITION
AQ472951.1 GI:4656517
GSS.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
AUTHORS
Zhao, S., Adams, M.D., Nierman, W., Malek, J., Shizuya, H., Simon, M. and
Venter, J.C.
Use of BAC End Sequences from Caltech Libraries for Sequence-Ready
Map Building
Unpublished (1997)
Other GSSs: CITBI-El-2589A16.TF
Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbe@tigr.org
Clones are available from Research Genetics (info@resgen.com). BAC
end search page:
http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html.
Seq primer: M13 Reverse
Class: BAC ends.

FEATURES
source
1. .337
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="2589A16"
/sex="male"
/cell_type="sperm"
/clone_lib="CITBI-El"
/note="Vector: pBelOBAC11; Site_1: EcoRI; Site_2: EcoRI;
Caltech Human BAC Library D"

ORIGIN
Query Match 81.7%; Score 18.8; DB 8; Length 337;
Best Local Similarity 90.9%; Pred. No. 4.5e+02;
Matches
20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY
1 CAGGAGATCCTGAGATTATGTG 22
|||||
Db
201 CAGGAGATCCTGAGATTATGTG 222
|||||

RESULT 11
AA608986
LOCUS
af05d07.s1 Soares testis NHT Homo sapiens cDNA clone IMAGE:1030765
3', similar to gb:J05158 CARBOXYPEPTIDASE N 83 KD CHAIN
(HUMAN); contains MER4.b2 MER4 MER4 repetitive element i, mRNA
sequence.
AA608986
DEFINITION
AA608986.1 GI:2457414
EST.
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 1 (bases 1 to 378)
 Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,
 Krizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M.,
 Martin, J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F.,
 Theising, B., White, Y., Wylie, T., Waterston, R. and Wilson, R.
 WashU-NCI human EST Project
 Unpublished (1997)
 Contact: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 This clone is available royalty-free through LNL; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 Insert Length: 533 Std Error: 0.00
 Seq primer: -40ml3 fwd. ET from Amersham
 High quality sequence stop: 367.
 Location/Qualifiers
 1..378
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:1030765"
 /sex="male"
 /lab_host="DH10B"
 /clone_lib="Soares testis NHT"
 /note="Vector: pT73D-Pac (Pharmacia) with a modified
 polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
 was prepared from mRNA obtained from Clontech
 Laboratories, Inc., and primed with a Not I - oligo(dT)
 primer [5'
 TGTATCCATCTGAAGTGGAGCGGCCCAATTTTTTTTTTTT 3']
 Double-stranded cDNA was ligated to Eco RI adaptors
 (Pharmacia), digested with Not I and cloned into the Not I
 and Eco RI sites of the modified pT73 vector. Library
 went through one round of normalization to Cot5, and was
 constructed by Bento Soares and M. Fatima Bonaldo."

Query Match 81.7%; Score 18.8; DB 1; Length 378;
 Best Local Similarity 90.9%; Pred. No. 4.6e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
 |||||
 Db 69 CAGGAGATCCTGAGAACATGTG 90

RESULT 12
 AQ279565/c
 LOCUS
 DEFINITION
 genomic survey sequence.
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 1 (bases 1 to 386)
 Berry, K., Granger, D., Suh, E., Wible, C., Shizuya, H., Simon, M. and
 Venter, J.C.
 Use of a random human BAC End Sequence Database for Sequence-Ready
 Map Building
 Unpublished (1998)
 Other GSSs: CITBI-E1-2523F6.TF
 Contact: Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research

9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: mdamas@tigr.org
 Clones are available from Research Genetics (info@resgen.com). BAC
 end search page:
 http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html.
 Seq primer: M13 Reverse
 Class: BAC ends.
 Location/Qualifiers
 1..386
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /clone="2523F6"
 /sex="male"
 /cell_type="sperm"
 /clone_lib="CITBI-E1"
 /note="Vector: pBelOBAC11; Site_1: EcoRI; Site_2: EcoRI;
 Caltech Human BAC Library D"

Query Match 81.7%; Score 18.8; DB 8; Length 386;
 Best Local Similarity 90.9%; Pred. No. 4.6e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
 |||||
 Db 155 CAGGAGATCCTGAGAACATGTG 134

RESULT 13
 AA569281
 LOCUS
 DEFINITION
 nm13e01.s1 NCI CGAP Lip2 Homo sapiens cDNA clone IMAGE:1061784
 similar to contains "Alu repetitive element; contains MER4.t3 MER4
 repetitive element ; mRNA sequence.
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 1 (bases 1 to 401)
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 Unpublished (1997)
 Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: L. Jeffrey Medeiros, M.D., Michael R.
 Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: David B. Krizman, Ph.D.
 cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 www-bio.llnl.gov/bbrp/image/image.html
 Insert Length: 684 Std Error: 0.00
 Seq primer: -40ml3 fwd. ET from Amersham
 High quality sequence stop: 359.
 Location/Qualifiers
 1..401
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:1061784"
 /tissue_type="liposarcoma"
 /lab_host="DH10B"
 /clone_lib="NCI CGAP Lip2"
 /note="Vector: pAMP10; mRNA made from liposarcoma, cDNA
 made by oligo-dT priming. Non-directionally cloned."

Size-selected on agarose gel, average insert size 600 bp.
Reference: Krizman et al. (1996) Cancer Research
56:5380-5383."

ORIGIN

Query Match 81.7%; Score 18.8; DB 1; Length 401;
Best Local Similarity 90.9%; Pred. No. 4.6e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||

Db 34 CAGGAGATCCTGAGAAATGTG 55
|||||

RESULT 14

AL703856

LOCUS 414 bp mRNA linear EST 04-SEP-2003
DEFINITION DKFZp686N0727_r1 686 (synonym: hlcc3) Homo sapiens cDNA clone
DKFZp686N0727 5', mRNA sequence.

ACCESSION

AL703856

VERSION AL703856.1 GI:19687211

KEYWORDS

EST.

SOURCE

Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 414)

Author: Ansgorge, W., Wirkner, U., Mewes, H.W., Weil, B. and Wiemann, S.

Title: EST (Ansgorge, W., Wirkner, U., Mewes, H.W., Weil, B. and Wiemann, S.)

Journal: Unpublished (1999)

Comment: Contact: MIPS

Features: MIPS

Ingolstaedter Landstr. 1, D-85764 Neuherberg, Germany

This is the 5' sequence of the clone insert

Clone from S. Wiemann, Molecular Genome Analysis, German Cancer

Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de;

sequenced by EMBL (European Molecular Biology Laboratories,

Heidelberg/Germany) within the cDNA sequencing consortium of the

German Genome Project.

No sl sequence available.

This clone (DKFZp686N0727) is available at the RZPD in Berlin.

Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059

Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.

Location/Qualifiers

1..414

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="DKFZp686N0727"

/dev_stage="adult"

/lab_host="DH10B"

/clone_lib="686 (synonym: hlcc3)"

/notes="Vector: pTriplex2; Site_1: SfIIA; Site_2: SfiIB;

cDNA-collection"

ORIGIN

Query Match 81.7%; Score 18.8; DB 1; Length 414;
Best Local Similarity 90.9%; Pred. No. 4.6e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
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Db 287 CAGGAGATCCTGAGACATGTG 308
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RESULT 15

AQ022522/c

LOCUS 441 bp DNA linear GSS 16-JUN-1998

DEFINITION HS 2180_A2_D01_MF CIT Approved Human Genomic Sperm Library D Homo

sapiens genomic clone Plate=2180 Col=2 Row=G, genomic survey

sequence.

ACCESSION AQ022522

VERSION AQ022522.1 GI:3220730

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

CONTACT

UNIVERSITY

TELEPHONE

FAX

EMAIL

SEQUENCE

PLATE

CLASS

HIGH QUALITY

SEQUENCE STOP

LOCATION/QUALIFIERS

1..441

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/clone="Plate=2180 Col=2 Row=G"

/sex="male"

/clone_lib="CIT Approved Human Genomic Sperm Library D"

/notes="Organ: sperm; Vector: pBelobAC11; BAC Clones in

E-Coli DH10B"

ORIGIN

Query Match 81.7%; Score 18.8; DB 8; Length 441;

Best Local Similarity 90.9%; Pred. No. 4.7e+02;

Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGAGATCCTGAGATTATGTG 22
|||||

Db 418 CAGGAGATCCTGAGACATGTG 397
|||||

Search completed: August 13, 2005, 06:45:12

Job time : 1485.12 secs

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